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【List of Submitted Items】

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【Name of item】	Abstract	1

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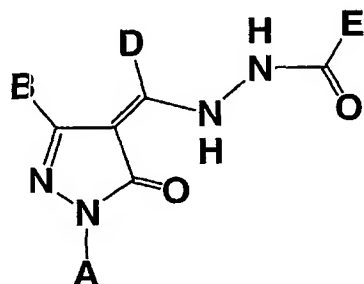
【Type of Document】

SCOPE OF THE CLAIM(S)

【Claim 1】

A pyrazolone compound represented by the formula (1)

【Ka 1】



(1)

5

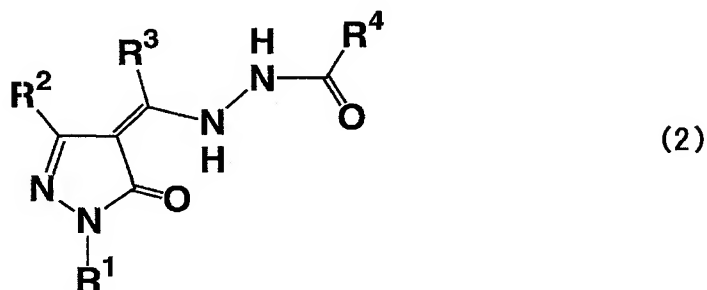
[wherein A is a C₂₋₁₄ aryl group (the C₂₋₁₄ aryl group may
 be optionally substituted with one or more C₁₋₆ alkyl
 groups, one or more C₁₋₃ alkyl groups substituted with one
 or more fluorine atoms, one or more halogen atoms, one or
 10 more nitro groups, one or more C₁₋₆ alkylcarbonyl groups,
 one or more hydroxyl groups or one or more amino groups
 (the hydroxyl groups and the amino groups may be
 substituted with a C₁₋₆ alkyl group or a C₁₋₆ alkylcarbonyl
 group)), B is a hydrogen atom, a C₁₋₆ alkyl group, a C₁₋₃
 15 alkyl group substituted with one or more fluorine atoms
 or a C₂₋₁₄ aryl group, D is a hydrogen atom, a C₁₋₆ alkyl
 group, a C₁₋₃ alkyl group substituted with one or more
 fluorine atoms or a C₂₋₁₄ aryl group, and E is a C₂₋₁₄ aryl
 group (the C₂₋₁₄ aryl group is optionally substituted with
 20 one or more hydroxyl groups, one or more nitro groups,
 one or more halogen atoms, NG¹G² (wherein G¹ and G² are
 independently hydrogen atoms, formyl groups, C₁₋₆ alkyl

groups or C₁₋₆ alkylcarbonyl groups), one or more carboxyl groups, one or more sulfonic acid groups, one or more phosphonic acid groups, one or more carbamido groups, one or more sulfamido groups, one or more hydroxycarbamido groups, one or more hydroxysulfamido groups, one or more tetrazole groups, one or more C₁₋₆ alkoxy carbonyl groups or X(CYZ)_nCO₂H (wherein X is CH₂, O, S or NG³ (G³ is a hydrogen atom, a C₁₋₆ alkyl group, a formyl group or a C₁₋₆ alkylcarbonyl group), Y and Z are independently hydrogen atoms or C₁₋₃ alkyl groups, and n is 0, 1, 2 or 3)), a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof.

【Claim 2】

A pyrazolone compound represented by the formula (2)

15 【Ka 2】



[wherein R¹ is a C₂₋₁₄ aryl group (the C₂₋₁₄ aryl group may be optionally substituted with one or more C₁₋₆ alkyl groups, one or more C₁₋₃ alkyl groups substituted with one or more fluorine atoms, one or more halogen atoms, one or more nitro groups, one or more C₁₋₆ alkylcarbonyl groups, one or more hydroxyl groups or one or more amino groups

(the hydroxyl groups and the amino groups may be substituted with a C₁₋₆ alkyl group or a C₁₋₆ alkylcarbonyl group)), R² is a hydrogen atom, a C₁₋₆ alkyl group, a C₁₋₃ alkyl group substituted with one or more fluorine atoms or a C₂₋₁₄ aryl group, R³ is a hydrogen atom, a C₁₋₆ alkyl group, a C₁₋₃ alkyl group substituted with one or more fluorine atoms or a C₂₋₁₄ aryl group, and R⁴ is a C₂₋₁₄ aryl group (the C₂₋₁₄ aryl group is optionally substituted with one or more hydroxyl groups, one or more nitro groups or NR⁵R⁶ (wherein R⁵ and R⁶ are independently hydrogen atoms, formyl groups, C₁₋₆ alkyl groups or C₁₋₆ alkylcarbonyl groups))], a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof.

[Claim 3]

The pyrazolone compound according to Claim 2, wherein R⁴ is a C₂₋₁₄ aryl group substituted with one or more hydroxyl groups, a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof.

[Claim 4]

The pyrazolone compound according to Claim 2, wherein R⁴ is a C₂₋₁₄ aryl group substituted with NR⁵R⁶ (wherein R⁵ and R⁶ are independently hydrogen atoms, formyl groups, C₁₋₆ alkyl groups or C₁₋₆ alkylcarbonyl groups), a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof.

[Claim 5]

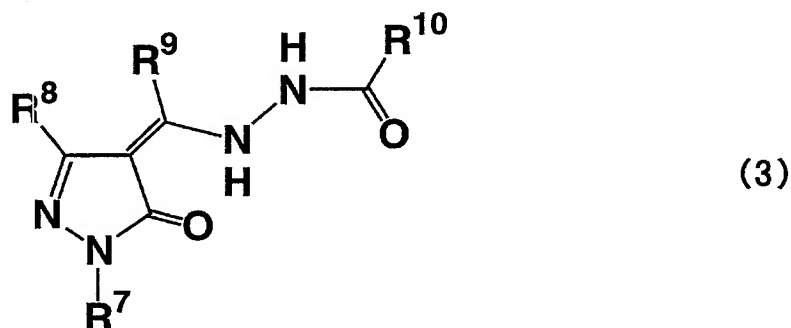
The pyrazolone compound according to Claim 2, wherein

R⁴ is a C₂₋₁₄ aryl group substituted with one or more nitro groups, a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof.

[Claim 6]

5 A pyrazolone compound represented by the formula (3)

[Ka 3.]



[wherein R⁷ is a C₂₋₁₄ aryl group (the C₂₋₁₄ aryl group may be optionally substituted with one or more C₁₋₆ alkyl groups, one or more C₁₋₃ alkyl groups substituted with one or more fluorine atoms, one or more halogen atoms, one or more nitro groups, one or more C₁₋₆ alkylcarbonyl groups, one or more hydroxyl groups or one or more amino groups (the hydroxyl groups and the amino groups may be substituted with a C₁₋₆ alkyl group or a C₁₋₆ alkylcarbonyl group)), R⁸ is a hydrogen atom, a C₁₋₆ alkyl group, a C₁₋₃ alkyl group substituted with one or more fluorine atoms or a C₂₋₁₄ aryl group, R⁹ is a hydrogen atom, a C₁₋₆ alkyl group, a C₁₋₃ alkyl group substituted with one or more fluorine atoms or a C₂₋₁₄ aryl group, and R¹⁰ is a C₂₋₁₄ aryl group (the C₂₋₁₄ aryl group is optionally substituted with one or more carboxyl groups, one or more sulfonic

acid groups, one or more phosphonic acid groups, one or more carbamido groups, one or more sulfamido groups, one or more hydroxycarbamido groups, one or more hydroxysulfamido groups, one or more tetrazole groups, one or more C₁₋₆ alkoxy carbonyl groups or X(CYZ)_nCO₂H (wherein X is CH₂, O, S or NR¹¹ (R¹¹ is a hydrogen atom, a C₁₋₆ alkyl group, a formyl group or a C₁₋₆ alkylcarbonyl group), Y and Z are independently hydrogen atoms or C₁₋₃ alkyl groups, and n is 0, 1, 2 or 3)), a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof.

【Claim 7】

The pyrazolone compound according to Claim 6, wherein R¹⁰ is a C₂₋₁₄ aryl group substituted with one or more carboxyl groups, a tautomer, prodrug or pharmaceutically acceptable salt of the compound, or a solvate thereof.

【Claim 8】

The pyrazolone compound according to Claim 6, wherein R¹⁰ is a C₂₋₁₄ aryl group substituted with X(CYZ)_nCO₂H (wherein X is CH₂, O, S or NR¹¹ (R¹¹ is a hydrogen atom, a C₁₋₆ alkyl group, a formyl group or a C₁₋₆ alkylcarbonyl group), Y and Z are independently hydrogen atoms or C₁₋₃ alkyl groups, and n is 0, 1, 2 or 3), a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof.

【Claim 9】

The pyrazolone compound according to Claim 6, wherein

R¹⁰ is a C₂₋₁₄ aryl group substituted with one or more sulfonic acid groups, a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof.

5 **【Claim 10】**

 The pyrazolone compound according to Claim 6, wherein R¹⁰ is a C₂₋₁₄ aryl group substituted with one or more phosphonic acid groups, a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a
10 solvate thereof.

【Claim 11】

 The pyrazolone compound according to Claim 6, wherein R¹⁰ is a C₂₋₁₄ aryl group substituted with one or more tetrazole groups, a tautomer, prodrug or pharmaceutically
15 acceptable salt of the compound or a solvate thereof.

【Claim 12】

 The pyrazolone compound according to Claim 6, wherein R¹⁰ is a C₂₋₁₄ aryl group substituted with one or more carbanido groups, a tautomer, prodrug or pharmaceutically
20 acceptable salt of the compound or a solvate thereof.

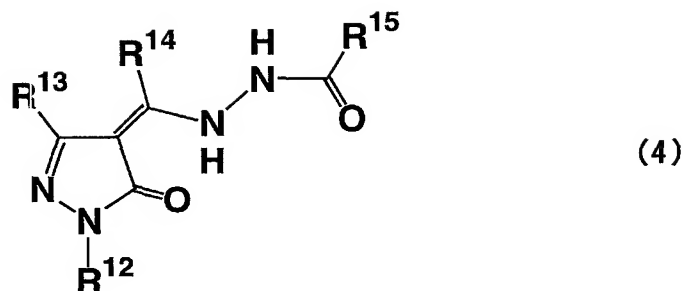
【Claim 13】

 The pyrazolone compound according to Claim 6, wherein R¹⁰ is a C₂₋₁₄ aryl group substituted with one or more sulfamido groups, a tautomer, prodrug or pharmaceutically
25 acceptable salt of the compound or a solvate thereof.

【Claim 14】

 A pyrazolone compound represented by the formula (4)

[Ka 4]



[wherein R¹² is a C₂₋₁₄ aryl group (the C₂₋₁₄ aryl group may be optionally substituted with one or more C₁₋₆ alkyl groups, one or more C₁₋₃ alkyl groups substituted with one or more fluorine atoms, one or more halogen atoms, one or more nitro groups, one or more C₁₋₆ alkylcarbonyl groups, one or more hydroxyl groups or one or more amino groups (the hydroxyl groups and the amino groups may be substituted with a C₁₋₆ alkyl group or a C₁₋₆ alkylcarbonyl group)), R¹³ is a hydrogen atom, a C₁₋₆ alkyl group, a C₁₋₃ alkyl group substituted with one or more fluorine atoms or a C₂₋₁₄ aryl group, R¹⁴ is a hydrogen atom, a C₁₋₆ alkyl group, a C₁₋₃ alkyl group substituted with one or more fluorine atoms or a C₂₋₁₄ aryl group, and R¹⁵ is a C₂₋₁₄ aryl group (the C₂₋₁₄ aryl group is substituted with a substituent selected from a hydroxyl group, an amino group, a nitro group and a halogen atom and with a substituent selected from a carboxyl group, a sulfonic acid group, a phosphonic acid group, a carbamido group, a sulfamido group, a hydroxycarbamido group, a hydroxysulfamido group, a tetrazole group, a C₁₋₆

alkoxycarbonyl group and $X(CYZ)_nCO_2H$ (wherein X is CH_2 , O, S or NR^{16} (R^{16} is a hydrogen atom, a C_{1-6} alkyl group, a formyl group or a C_{1-6} alkylcarbonyl group), Y and Z are independently hydrogen atoms or C_{1-3} alkyl groups, and n is 0, 1, 2 or 3)), a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof.

【Claim 15】

The pyrazolone compound according to Claim 14, wherein R^{15} is a C_{2-14} aryl group substituted with a hydroxyl group and a carboxyl group, a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof.

【Claim 16】

The pyrazolone compound according to Claim 14, wherein R^{15} is a C_{2-14} aryl group substituted with an amino group and a carboxyl group, a tautomer, a prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof.

【Claim 17】

The pyrazolone compound according to Claim 14, wherein R^{15} is a C_{2-14} aryl group substituted with a substituent selected from a nitro group and a halogen atom and with a carboxyl group, a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof.

【Claim 18】

The thrombopoietin receptor activator according to Claim 1.

【Claim 19】

5 The thrombopoietin receptor activator according to Claim 2.

【Claim 20】

The thrombopoietin receptor activator according to Claim 3.

10 **【Claim 21】**

The thrombopoietin receptor activator according to Claim 4.

【Claim 22】

15 The thrombopoietin receptor activator according to Claim 5.

【Claim 23】

The thrombopoietin receptor activator according to Claim 6.

【Claim 24】

20 The thrombopoietin receptor activator according to Claim 7.

【Claim 25】

The thrombopoietin receptor activator according to Claim 8.

25 **【Claim 26】**

The thrombopoietin receptor activator according to Claim 9.

【Claim 27】

The thrombopoietin receptor activator according to Claim 10.

【Claim 28】

5 The thrombopoietin receptor activator according to Claim 11.

【Claim 29】

The thrombopoietin receptor activator according to Claim 12.

10 **【Claim 30】**

The thrombopoietin receptor activator according to Claim 13.

【Claim 31】

15 The thrombopoietin receptor activator according to Claim 14.

【Claim 32】

The thrombopoietin receptor activator according to Claim 15.

【Claim 33】

20 The thrombopoietin receptor activator according to Claim 16.

【Claim 34】

The thrombopoietin receptor activator according to Claim 17.

25 **【Claim 35】**

A preventive, therapeutic or improving agent for diseases against which activation of the thrombopoietin

receptor is effective, which contains the thrombopoietin
receptor activator according to Claim 18, Claim 19, Claim
20, Claim 21, Claim 22, Claim 23, Claim 24, Claim 25,
Claim 26, Claim 27, Claim 28, Claim 29, Claim 30, Claim
5 31, Claim 32, Claim 33 or Claim 34, a tautomer, prodrug
or pharmaceutically acceptable salt of the activator or a
solvate thereof, as an active ingredient.

【Claim 36】

A platelet increasing agent containing the
10 thrombopoietin receptor activator according to Claim 18,
Claim 19, Claim 20, Claim 21, Claim 22, Claim 23, Claim
24, Claim 25, Claim 26, Claim 27, Claim 28, Claim 29,
Claim 30, Claim 31, Claim 32, Claim 33 or Claim 34, a
tautomer, prodrug or pharmaceutically acceptable salt of
15 the activator or a solvate thereof, as an active
ingredient.

【Type of Document】 DESCRIPTION

【Title of the invention】

PYRAZOLONE COMPOUNDS AND THROMBOPOIETIN RECEPTOR
ACTIVATOR

5 【Technical field】

The present invention relates to preventive,
therapeutic and improving agents having affinity for and
agonistic action on the thrombopoietin receptor for
diseases against which activation of the thrombopoietin
10 receptor is effective. Specifically, it relates to
pharmaceutical compositions comprising compounds which
increase platelets through stimulation of differentiation
and proliferation of hematopoietic stem cells,
megakaryocytic progenitor cells and megakaryocytes or
15 compounds for therapeutic angiogenesis or with anti-
arteriosclerosis action that stimulate differentiation
and proliferation of vascular endothelial cells and
endothelial progenitor cells.

【Background art】

20 Thrombopoietin is a cytokine consisting of 332 amino
acids that increases platelet production by stimulating
differentiation and proliferation of hematopoietic stem
cells, megakaryocytic progenitor cells and megakaryocytes
mediated by its receptor and therefore is promising as a
25 drug for hematological disorders. Recent reports that it
stimulates differentiation and proliferation of vascular
endothelial cells and endothelial progenitor cells have

raised expectations of therapeutic angiogenesis, anti-arteriosclerosis and prevention of cardiovascular events (for example, non-patent document 1, non-patent document 2 and non-patent document 3).

5 Biologically active substances which have been known so far to regulate platelet production through the thrombopoietin receptor include, in addition to thrombopoietin itself, low molecular weight peptides having affinity for the thrombopoietin receptor (for
10 example, patent document 1, patent document 2, patent document 3 and patent document 4).

As a result of search for nonpeptidic low molecular weight compounds that increase platelet production mediated by the thrombopoietin receptor, low molecular
15 weight compounds having affinity for the thrombopoietin receptor have been reported (for example, patent document 5 to patent document 22).

- 1) Applications filed by Hokuriku Seiyaku Co., Ltd. relating to 1,4-benzodiazepine derivatives (patent
20 documents 5 and 6)
- 2) International Laid-open Patent Applications filed by Shionogi & Co., Ltd. (patent documents 7-10)
- 3) International Laid-open Patent Applications filed by SmithKline Beecham Corp (patent documents 11-19)
- 25 4) Japanese Laid-open Patent Application filed by Torii Pharmaceutical Co., Ltd. (patent document 20)
- 5) International Laid-open Patent Application filed by

Roche Diagnostics GMBH (patent document 21)

6) International Laid-open Patent Application filed by
Yamanouchi Pharmaceutical Co., Ltd. (patent document 22)

Some reports have been made about pyrazolone
5 compounds (such as non-patent documents 4-13).

- 【Patent Document 1】 JP-A-10-72492
- 【Patent Document 2】 WO96/40750
- 【Patent Document 3】 WO96/40189
- 【Patent Document 4】 WO98/25965
- 10 【Patent Document 5】 JP-A-11-1477
- 【Patent Document 6】 JP-A-11-152276
- 【Patent Document 7】 WO01/07423
- 【Patent Document 8】 WO01/53267
- 【Patent Document 9】 WO02/059099
- 15 【Patent Document 10】 WO02/059100
- 【Patent Document 11】 WO00/35446
- 【Patent Document 12】 WO00/66112
- 【Patent Document 13】 WO01/34585
- 【Patent Document 14】 WO01/17349
- 20 【Patent Document 15】 WO01/39773
- 【Patent Document 16】 WO01/21180
- 【Patent Document 17】 WO01/89457
- 【Patent Document 18】 WO02/49413
- 【Patent Document 19】 WO02/085343
- 25 【Patent Document 20】 JP-A-2001-97948
- 【Patent Document 21】 WO99/11262
- 【Patent Document 22】 WO02/062775

【Non-patent Document 1】 Microvasc Res 1999: 58,
p.108-113

【Non-patent Document 2】 Circ Res 1999: 84, p.785-796

【Non-patent Document 3】 Blood 2001:98, p.71a

5 【Non-patent Document 4】 Huaxue Xuebao (2001), 59(9)
p.1495-1501

【Non-patent Document 5】 Synthesis and Reactivity in
Inorganic andMetal Organic Chemistry (2000), 30(7)
p.1265-1271

10 【Non-patent Document 6】 Synthesis and Reactivity in
Inorganic andMetal Organic Chemistry (2002), 32(4) p.739-
751

 【Non-patent Document 7】 Synthesis and Reactivity in
Inorganic andMetal Organic Chemistry (2002), 32(5) p.903-
15 912

 【Non-patent Document 8】 Jiegou Huaxue (2002), 21(5),
p.553-556

 【Non-patent Document 9】 Polyhedroon (1997), 16(11)
p.1825-1829

20 【Non-patent Document 10】 Arzneim-Forsch (1969),
19(10) p.1721-1723

 【Non-patent Document 11】 Structural Chemistry (1999),
10(2), 105-119

 【Non-patent Document 12】 Chemical Sciences (1996),
25 51(9), 1240-1244

 【Non-patent Document 13】 Chemical Sciences (1997),
52(2), 237-242

【Disclosure of Invention】

【Problems to be solved by the invention】

Thrombopoietin and low molecular weight peptides having affinity for the thrombopoietin receptor are likely to be easily degraded in the gastrointestinal tract and are usually difficult to orally administer. As to thrombopoietin itself, the appearance of anti-thrombopoietin antibodies have been reported.

Besides, though it is probably possible to orally administer nonpeptidic low molecular weight compounds, no practical drugs have been put on the market.

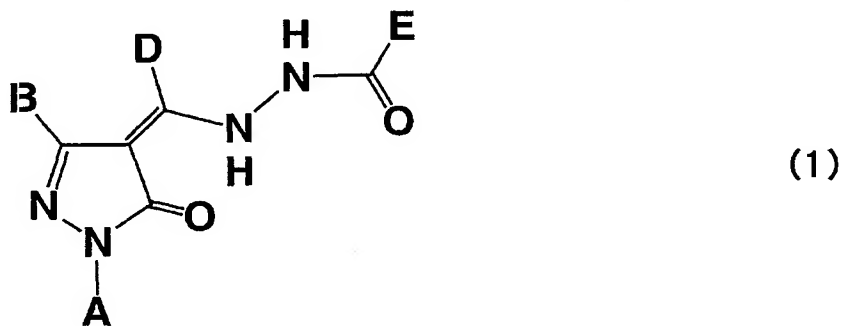
Therefore, orally administrable low molecular weight compounds having excellent affinity for and agonistic action on the thrombopoietin receptor as preventive, therapeutic and improving agents for diseases against which activation of the thrombopoietin receptor is effective have been demanded. Specifically, low molecular weight compounds which can serve as platelet increasing agents or increasing agents for other blood cells by stimulating differentiation and proliferation of hematopoietic stem cells, megakaryocytic progenitor cells and megakaryocytes or low molecular weight compounds which can be used for therapeutic angiogenesis or as preventive and therapeutic agents for arteriosclerosis by stimulating endothelial cells and endothelial progenitor cells have been demanded.

[Means for solving problem]

The present inventors conducted extensive research to find low molecular weight compounds having affinity for and agonistic action on the thrombopoietin receptor, and
5 as a result, found that the compounds of the present invention have high affinity and agonistic action which enable them to show potent platelet increasing action by stimulating differentiation and proliferation of megakaryocytic progenitor cells and megakaryocytes. The
10 present invention was accomplished on the basis of this discovery.

Namely, the present invention relates to a pyrazolone compound represented by the formula (1)

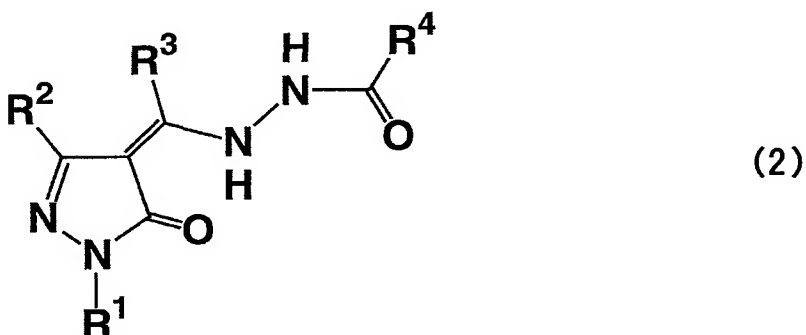
[Ka 5]



15 [wherein A is a C₂₋₁₄ aryl group (the C₂₋₁₄ aryl group may be optionally substituted with one or more C₁₋₆ alkyl groups, one or more C₁₋₃ alkyl groups substituted with one or more fluorine atoms, one or more halogen atoms, one or
20 more nitro groups, one or more C₁₋₆ alkylcarbonyl groups, one or more hydroxyl groups or one or more amino groups (the hydroxyl groups and the amino groups may be

substituted with a C₁₋₆ alkyl group or a C₁₋₆ alkylcarbonyl group)), B is a hydrogen atom, a C₁₋₆ alkyl group, a C₁₋₃ alkyl group substituted with one or more fluorine atoms or a C₂₋₁₄ aryl group, D is a hydrogen atom, a C₁₋₆ alkyl group, a C₁₋₃ alkyl group substituted with one or more fluorine atoms or a C₂₋₁₄ aryl group, and E is a C₂₋₁₄ aryl group (the C₂₋₁₄ aryl group is optionally substituted with one or more hydroxyl groups, one or more nitro groups, one or more halogen atoms, NG¹G² (wherein G¹ and G² are independently hydrogen atoms, formyl groups, C₁₋₆ alkyl groups or C₁₋₆ alkylcarbonyl groups), one or more carboxyl groups, one or more sulfonic acid groups, one or more phosphonic acid groups, one or more carbamido groups, one or more sulfamido groups, one or more hydroxycarbamido groups, one or more hydroxysulfamido groups, one or more tetrazole groups, one or more C₁₋₆ alkoxy carbonyl groups or X(CYZ)_nCO₂H (wherein X is CH₂, O, S or NG³ (G³ is a hydrogen atom, a C₁₋₆ alkyl group, a formyl group or a C₁₋₆ alkylcarbonyl group), Y and Z are independently hydrogen atoms or C₁₋₃ alkyl groups, and n is 0, 1, 2 or 3))), a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof, a thrombopoietin receptor activator, a preventive, therapeutic or improving agent for diseases against which activation of the thrombopoietin receptor is effective which contains the thrombopoietin receptor activator, a tautomer, prodrug or pharmaceutically acceptable salt of the

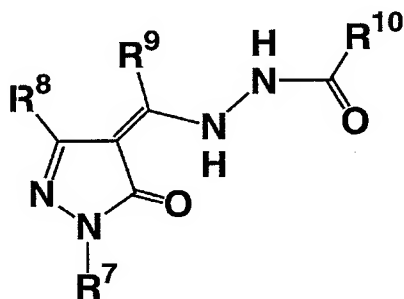
thrombopoietin receptor activator or a solvate thereof as
an active ingredient, and a platelet increasing agent
containing the thrombopoietin receptor activator, a
tautomer, prodrug or pharmaceutically acceptable salt of
5 the thrombopoietin receptor activator or a solvate
thereof as an active ingredient. It also relates to a
pyrazolone compound represented by the formula (2)
[Ka 6]



10 [wherein R¹ is a C₂₋₁₄ aryl group (the C₂₋₁₄ aryl group may
be optionally substituted with one or more C₁₋₆ alkyl
groups, one or more C₁₋₃ alkyl groups substituted with one
or more fluorine atoms, one or more halogen atoms, one or
more nitro groups, one or more C₁₋₆ alkylcarbonyl groups,
15 one or more hydroxyl groups or one or more amino groups
(the hydroxyl groups and the amino groups may be
substituted with a C₁₋₆ alkyl group or a C₁₋₆ alkylcarbonyl
group)), R² is a hydrogen atom, a C₁₋₆ alkyl group, a C₁₋₃
alkyl group substituted with one or more fluorine atoms
20 or a C₂₋₁₄ aryl group, R³ is a hydrogen atom, a C₁₋₆ alkyl
group, a C₁₋₃ alkyl group substituted with one or more

fluorine atoms or a C₂₋₁₄ aryl group, and R⁴ is a C₂₋₁₄ aryl group (the C₂₋₁₄ aryl group is optionally substituted with one or more hydroxyl groups, one or more nitro groups or NR⁵R⁶ (wherein R⁵ and R⁶ are independently hydrogen atoms, formyl groups, C₁₋₆ alkyl groups or C₁₋₆ alkylcarbonyl groups))], a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof, a thrombopoietin receptor activator, a preventive, therapeutic or improving agent for diseases against which activation of the thrombopoietin receptor is effective which contains the thrombopoietin receptor activator, a tautomer, prodrug or pharmaceutically acceptable salt of the thrombopoietin receptor activator or a solvate thereof as an active ingredient, and a platelet increasing agent containing the thrombopoietin receptor activator, a tautomer, prodrug or pharmaceutically acceptable salt of the thrombopoietin receptor activator or a solvate thereof as an active ingredient. It further relates to a pyrazolone compound represented by the formula (3)

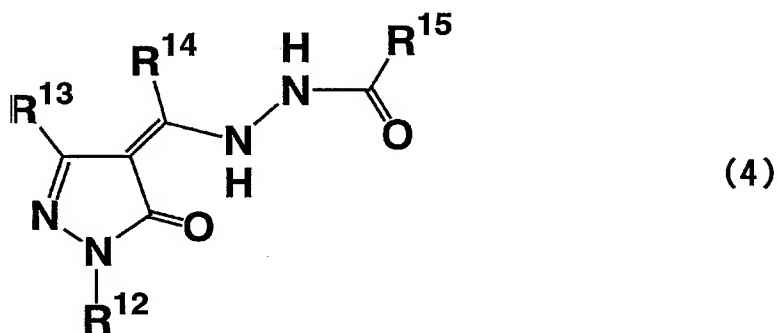
【Ka 7】



(3)

[wherein R^7 is a C_{2-14} aryl group (the C_{2-14} aryl group may be optionally substituted with one or more C_{1-6} alkyl groups, one or more C_{1-3} alkyl groups substituted with one or more fluorine atoms, one or more halogen atoms, one or more nitro groups, one or more C_{1-6} alkylcarbonyl groups, one or more hydroxyl groups or one or more amino groups (the hydroxyl groups and the amino groups may be substituted with a C_{1-6} alkyl group or a C_{1-6} alkylcarbonyl group)), R^8 is a hydrogen atom, a C_{1-6} alkyl group, a C_{1-3} alkyl group substituted with one or more fluorine atoms or a C_{2-14} aryl group, R^9 is a hydrogen atom, a C_{1-6} alkyl group, a C_{1-3} alkyl group substituted with one or more fluorine atoms or a C_{2-14} aryl group, and R^{10} is a C_{2-14} aryl group (the C_{2-14} aryl group is optionally substituted with one or more carboxyl groups, one or more sulfonic acid groups, one or more phosphonic acid groups, one or more carbamido groups, one or more sulfamido groups, one or more hydroxycarbamido groups, one or more hydroxysulfamido groups, one or more tetrazole groups, one or more C_{1-6} alkoxy carbonyl groups or $X(CYZ)_nCO_2H$ (wherein X is CH_2 , O , S or NR^{11} (R^{11} is a hydrogen atom, a C_{1-6} alkyl group, a formyl group or a C_{1-6} alkylcarbonyl group), Y and Z are independently hydrogen atoms or C_{1-3} alkyl groups, and n is 0, 1, 2 or 3))], a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof, a thrombopoietin receptor activator, a preventive, therapeutic or improving agent

for diseases against which activation of the
thrombopoietin receptor is effective which contains the
thrombopoietin receptor activator, a tautomer, prodrug or
pharmaceutically acceptable salt of the thrombopoietin
5 receptor activator or a solvate thereof as an active
ingredient, and a platelet increasing agent containing
the thrombopoietin receptor activator, a tautomer,
prodrug or pharmaceutically acceptable salt of the
thrombopoietin receptor activator or a solvate thereof as
10 an active ingredient. It still further relates to a
pyrazolone compound represented by the formula (4)
[Ka 8]



[wherein R^{12} is a C_{2-14} aryl group (the C_{2-14} aryl group may
15 be optionally substituted with one or more C_{1-6} alkyl
groups, one or more C_{1-3} alkyl groups substituted with one
or more fluorine atoms, one or more halogen atoms, one or
more nitro groups, one or more C_{1-6} alkylcarbonyl groups,
one or more hydroxyl groups or one or more amino groups
20 (the hydroxyl groups and the amino groups may be
substituted with a C_{1-6} alkyl group or a C_{1-6} alkylcarbonyl

group)), R^{13} is a hydrogen atom, a C_{1-6} alkyl group, a C_{1-3} alkyl group substituted with one or more fluorine atoms or a C_{2-14} aryl group, R^{14} is a hydrogen atom, a C_{1-6} alkyl group, a C_{1-3} alkyl group substituted with one or more
5 fluorine atoms or a C_{2-14} aryl group, and R^{15} is a C_{2-14} aryl group (the C_{2-14} aryl group is substituted with a substituent selected from a hydroxyl group, an amino group, a nitro group and a halogen atom and with a substituent selected from a carboxyl group, a sulfonic
10 acid group, a phosphonic acid group, a carbamido group, a sulfamido group, a hydroxycarbamido group, a hydroxysulfamido group, a tetrazole group, a C_{1-6} alkoxy carbonyl group and $X(CYZ)_nCO_2H$ (wherein X is CH_2 , O, S or NR^{16} (R^{16} is a hydrogen atom, a C_{1-6} alkyl group, a
15 formyl group or a C_{1-6} alkyl carbonyl group), Y and Z are independently hydrogen atoms or C_{1-3} alkyl groups, and n is 0, 1, 2 or 3))), a tautomer, prodrug or pharmaceutically acceptable salt of the compound or a solvate thereof, a thrombopoietin receptor activator, a
20 preventive, therapeutic or improving agent for diseases against which activation of the thrombopoietin receptor is effective which contains the thrombopoietin receptor activator, a tautomer, prodrug or pharmaceutically acceptable salt of the thrombopoietin receptor activator
25 or a solvate thereof as an active ingredient, and a platelet increasing agent containing the thrombopoietin receptor activator, a tautomer, prodrug or

pharmaceutically acceptable salt of the thrombopoietin receptor activator or a solvate thereof as an active ingredient.

Though WO99/11262 (patent document 21), WO01/34585
5 (patent document 13), WO02/49413 (patent document 18) disclose pyrazolone compounds having platelet increasing action, there is no specific disclosure of the pyrazolone compounds of the present invention. The compounds of the present invention showed high activity that could not be
10 expected from the disclosure in WO99/11262 (patent document 21), WO01/34585 (patent document 13) or WO02/49413 (patent document 18).

【Best mode(s) for carrying out the invention】

Now, the present invention will be described in
15 detail.

In the present invention, "n" denotes normal, "i" denotes iso, "s" denotes secondary, "t" denotes tertiary, "c" denotes cyclo, "o" denotes ortho, "m" denotes meta, "p" denotes para, "Ph" denotes phenyl, "Py" denotes
20 pyridyl, "Naphthyl" denotes naphthyl, "Me" denotes methyl, "Et" denotes ethyl, "Pr" denotes propyl, and "Bu" denotes butyl.

First, the terms in the respective substituents A, B, D, E, G¹, G², G³, R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰
25 R¹¹, R¹², R¹³, R¹⁴, R¹⁵ and R¹⁶ will be explained.

As a halogen atom, fluorine, chlorine, bromine or iodine may be mentioned.

A C₁₋₃ alkyl group may be linear, branched or a C₃ cycloalkyl group, and methyl, ethyl, n-propyl, i-propyl and c-propyl and the like may be mentioned. A C₁₋₆ alkyl group may be linear, branched or a C₃₋₆ cycloalkyl group, and in addition to those mentioned above, n-butyl, i-butyl, s-butyl, t-butyl, c-butyl, 1-methyl-c-propyl, 2-methyl-c-propyl, n-pentyl, 1-methyl-n-butyl, 2-methyl-n-butyl, 3-methyl-n-butyl, 1,1-dimethyl-n-propyl, 1,2-dimethyl-n-propyl, 2,2-dimethyl-n-propyl, 1-ethyl-n-propyl, c-pentyl, 1-methyl-c-butyl, 2-methyl-c-butyl, 3-methyl-c-butyl, 1,2-dimethyl-c-propyl, 2,3-dimethyl-c-propyl, 1-ethyl-c-propyl, 2-ethyl-c-propyl, n-hexyl, 1-methyl-n-pentyl, 2-methyl-n-pentyl, 3-methyl-n-pentyl, 4-methyl-n-pentyl, 1,1-dimethyl-n-butyl, 1,2-dimethyl-n-butyl, 1,3-dimethyl-n-butyl, 2,2-dimethyl-n-butyl, 2,3-dimethyl-n-butyl, 3,3-dimethyl-n-butyl, 1-ethyl-n-butyl, 2-ethyl-n-butyl, 1,1,2-trimethyl-n-propyl, 1,2,2-trimethyl-n-propyl, 1-ethyl-1-methyl-n-propyl, 1-ethyl-2-methyl-n-propyl, c-hexyl, 1-methyl-c-pentyl, 2-methyl-c-pentyl, 3-methyl-c-pentyl, 1-ethyl-c-butyl, 2-ethyl-c-butyl, 3-ethyl-c-butyl, 1,2-dimethyl-c-butyl, 1,3-dimethyl-c-butyl, 2,2-dimethyl-c-butyl, 2,3-dimethyl-c-butyl, 2,4-dimethyl-c-butyl, 3,3-dimethyl-c-butyl, 1-n-propyl-c-propyl, 2-n-propyl-c-propyl, 1-i-propyl-c-propyl, 2-i-propyl-c-propyl, 1,2,2-trimethyl-c-propyl, 1,2,3-trimethyl-c-propyl, 2,2,3-trimethyl-c-propyl, 1-ethyl-2-methyl-c-propyl, 2-ethyl-1-methyl-c-propyl, 2-

ethyl-2-methyl-c-propyl, 2-ethyl-3-methyl-c-propyl and the like may be mentioned.

A C₂₋₁₄ aryl group may be a C₆₋₁₄ aryl group containing no hetero atoms as ring constituting atoms or a C₂₋₉ aromatic heterocyclic group, and a C₂₋₉ aromatic heterocyclic group may be a 5 to 7-membered C₂₋₆ heteromonocyclic group or 8 to 10-membered C₅₋₉ fused heterobicyclic group containing from 1 to 3 oxygen atoms, nitrogen atoms or sulfur atoms singly or in combination.

10 As a C₆₋₁₄ aryl group containing no hetero atoms, a phenyl group, a 1-indenyl group, a 2-indenyl group, a 3-indenyl group, a 4-indenyl group, a 5-indenyl group, a 6-indenyl group, a 7-indenyl group, an α -naphthyl group, a β -naphthyl group, a 1-tetrahydronaphthyl group, a 2-tetrahydronaphthyl group, a 5-tetrahydronaphthyl group, a 6-tetrahydronaphthyl group, an o-biphenyl group, a m-biphenyl group, a p-biphenyl group, a 1-anthryl group, a 2-anthryl group, a 9-anthryl group, a 1-phenanthryl group, a 2-phenanthryl group, a 3-phenanthryl group, a 4-phenanthryl group, a 9-phenanthryl group or the like may be mentioned.

25 A 5 to 7-membered C₂₋₆ heteromonocyclic group may be a 2-thienyl group, a 3-thienyl group, a 2-furyl group, a 3-furyl group, a 2-pyranyl group, a 3-pyranyl group, a 4-pyranyl group, a 1-pyrrolyl group, a 2-pyrrolyl group, a 3-pyrrolyl group, a 1-imidazolyl group, a 2-imidazolyl group, a 4-imidazolyl group, a 1-pyrazolyl group, a 3-

pyrazolyl group, a 4-pyrazolyl group, a 2-thiazolyl group, a 4-thiazolyl group, a 5-thiazolyl group, a 3-isothiazolyl group, a 4-isothiazolyl group, a 5-isothiazolyl group, a 2-oxazolyl group, a 4-oxazolyl group, a 5-oxazolyl group, a 3-isoxazolyl group, a 4-isoxazolyl group, a 5-isoxazolyl group, a 2-pyridyl group, a 3-pyridyl group, a 4-pyridyl group, a 2-pyrazinyl group, a 2-pyrimidinyl group, a 4-pyrimidinyl group, a 5-pyrimidinyl group, a 3-pyridazinyl group, a 4-pyridazinyl group, a 2-1,3,4-oxadiazolyl group, a 2-1,3,4-thiadiazolyl group, a 3-1,2,4-oxadiazolyl group, a 5-1,2,4-oxadiazolyl group, a 3-1,2,4-thiadiazolyl group, a 5-1,2,4-thiadiazolyl group, a 3-1,2,5-oxadiazolyl group, a 3-1,2,5-thiadiazolyl group or the like.

15 A 8 to 10-membered C₅₋₉ fused heterocyclic group may be a 2-benzofuranyl group, a 3-benzofuranyl group, a 4-benzofuranyl group, a 5-benzofuranyl group, a 6-benzofuranyl group, a 7-benzofuranyl group, a 1-isobenzofuranyl group, a 4-isobenzofuranyl group, a 5-isobenzofuranyl group, a 2-benzothienyl group, a 3-benzothienyl group, a 4-benzothienyl group, a 5-benzothienyl group, a 6-benzothienyl group, a 7-benzothienyl group, a 1-isobenzothienyl group, a 4-isobenzothienyl group, a 5-isobenzothienyl group, a 2-chromenyl group, a 3-chromenyl group, a 4-chromenyl group, a 5-chromenyl group, a 6-chromenyl group, a 7-chromenyl group, a 8-chromenyl group, a 1-indolizinyl

group, a 2-indoliziny1 group, a 3-indoliziny1 group, a 5-indoliziny1 group, a 6-indoliziny1 group, a 7-indoliziny1 group, a 8-indoliziny1 group, a 1-isoindoly1 group, a 2-isoindoly1 group, a 4-isoindoly1 group, a 5-isoindoly1
5 group, a 1-indoly1 group, a 2-indoly1 group, a 3-indoly1 group, a 4-indoly1 group, a 5-indoly1 group, a 6-indoly1 group, a 7-indoly1 group, 1-indazolyl group, a 2-indazolyl group, a 3-indazolyl group, a 4-indazolyl group, a 5-indazolyl group, a 6-indazolyl group, a 7-indazolyl group, a 1-puriny1 group, a 2-puriny1 group, a
10 3-puriny1 group, a 6-puriny1 group, a 7-puriny1 group, a 8-puriny1 group, a 2-quinoly1 group, a 3-quinoly1 group, a 4-quinoly1 group, a 5-quinoly1 group, a 6-quinoly1 group, a 7-quinoly1 group, a 8-quinoly1 group, a 1-isoquinoly1 group, a 3-isoquinoly1 group, a 4-isoquinoly1 group, a 5-isoquinoly1 group, a 6-isoquinoly1 group, a 7-isoquinoly1 group, a 8-isoquinoly1 group, a 1-phthalaziny1 group, a 5-phthalaziny1 group, a 6-phthalaziny1 group, a 1-2,7-naphthyridiny1 group, a 3-
20 2,7-naphthyridiny1 group, a 4-2,7-naphthyridiny1 group, a 1-2,6-naphthyridiny1 group, a 3-2,6-naphthyridiny1 group, a 4-2,6-naphthyridiny1 group, a 2-1,8-naphthyridiny1 group, a 3-1,8-naphthyridiny1 group, a 4-1,8-naphthyridiny1 group, a 2-1,7-naphthyridiny1 group, a 3-1,7-naphthyridiny1 group, a 4-1,7-naphthyridiny1 group, a
25 5-1,7-naphthyridiny1 group, a 6-1,7-naphthyridiny1 group, a 8-1,7-naphthyridiny1 group, 2-1,6-naphthyridiny1 group,

a 3-1,6-naphthyridinyl group, a 4-1,6-naphthyridinyl group, a 5-1,6-naphthyridinyl group, a 7-1,6-naphthyridinyl group, a 8-1,6-naphthyridinyl group, a 2-1,5-naphthyridinyl group, a 3-1,5-naphthyridinyl group, a 4-1,5-naphthyridinyl group, a 6-1,5-naphthyridinyl group, a 7-1,5-naphthyridinyl group, a 8-1,5-naphthyridinyl group, a 2-quinoxalinylnyl group, a 5-quinoxalinylnyl group, a 6-quinoxalinylnyl group, a 2-quinazolinyl group, a 4-quinazolinyl group, a 5-quinazolinyl group, a 6-quinazolinyl group, a 7-quinazolinyl group, a 8-quinazolinyl group, a 3-cinnolinyl group, a 4-cinnolinyl group, a 5-cinnolinyl group, a 6-cinnolinyl group, a 7-cinnolinyl group, a 8-cinnolinyl group, a 2-pteridinyl group, a 4-pteridinyl group, a 6-pteridinyl group, a 7-pteridinyl group or the like.

A C₁₋₃ alkyl group substituted with one or more fluorine atoms may be a trifluoromethyl group, a difluoromethyl group, a monofluoromethyl group, a pentafluoroethyl group, a 1,1-difluoro-2,2-difluoroethyl group, a heptafluoropropyl group or the like.

A C₁₋₆ alkylcarbonyl group may be methylcarbonyl, ethylcarbonyl, n-propylcarbonyl, i-propylcarbonyl, n-butylcarbonyl, i-butylcarbonyl, s-butylcarbonyl, t-butylcarbonyl, n-pentylcarbonyl, 1-methyl-n-butylcarbonyl, 2-methyl-n-butylcarbonyl, 3-methyl-n-butylcarbonyl, 1,1-dimethyl-n-propylcarbonyl, 1,2-dimethyl-n-propylcarbonyl, 2,2-dimethyl-n-propylcarbonyl,

1-ethyl-n-propylcarbonyl, n-hexylcarbonyl, 1-methyl-n-pentylcarbonyl, 2-methyl-n-pentylcarbonyl, 3-methyl-n-pentylcarbonyl, 4-methyl-n-pentylcarbonyl, 1,1-dimethyl-n-butylcarbonyl, 1,2-dimethyl-n-butylcarbonyl, 1,3-
5 dimethyl-n-butylcarbonyl, 2,2-dimethyl-n-butylcarbonyl, 2,3-dimethyl-n-butylcarbonyl, 3,3-dimethyl-n-butylcarbonyl, 1-ethyl-n-butylcarbonyl, 2-ethyl-n-butylcarbonyl, 1,1,2-trimethyl-n-propylcarbonyl, 1,2,2-trimethyl-n-propylcarbonyl, 1-ethyl-1-methyl-n-
10 propylcarbonyl, 1-ethyl-2-methyl-n-propylcarbonyl or the like.

A C₁₋₆ alkoxy carbonyl group may be methoxycarbonyl, ethoxycarbonyl, n-propoxycarbonyl, i-propoxycarbonyl, n-butoxycarbonyl, i-butoxycarbonyl, s-butoxycarbonyl, t-
15 butoxycarbonyl, n-pentyloxycarbonyl, 1-methyl-n-butoxycarbonyl, 2-methyl-n-butoxycarbonyl, 3-methyl-n-butoxycarbonyl, 1,1-dimethyl-n-propoxycarbonyl, 1,2-dimethyl-n-propoxycarbonyl, 2,2-dimethyl-n-propoxycarbonyl, 1-ethyl-n-propoxycarbonyl, n-
20 hexyloxycarbonyl, 1-methyl-n-pentyloxycarbonyl, 2-methyl-n-pentyloxycarbonyl, 3-methyl-n-pentyloxycarbonyl, 4-methyl-n-pentyloxycarbonyl, 1,1-dimethyl-n-butoxycarbonyl, 1,2-dimethyl-n-butoxycarbonyl, 1,3-dimethyl-n-butoxycarbonyl, 2,2-dimethyl-n-butoxycarbonyl,
25 2,3-dimethyl-n-butoxycarbonyl, 3,3-dimethyl-n-butoxycarbonyl, 1-ethyl-n-butoxycarbonyl, 2-ethyl-n-butoxycarbonyl, 1,1,2-trimethyl-n-propoxycarbonyl, 1,2,2-

trimethyl-n-propoxycarbonyl, 1-ethyl-1-methyl-n-propoxycarbonyl, 1-ethyl-2-methyl-n-propoxycarbonyl or the like.

Specific preferred examples of the substituents A,
5 R¹, R⁷ and R¹² are a phenyl group, thienyl groups (a 2-thienyl group and a 3-thienyl group), furyl groups (a 2-furyl group and a 3-furyl group), pyridazinyl groups (a 3-pyridazinyl group and a 4-pyridazinyl group), pyridyl groups (a 2-pyridyl group, a 3-pyridyl group and a 4-
10 pyridyl group), quinolyl groups (a 2-quinolyl group, a 3-quinolyl group, a 4-quinolyl group, a 5-quinolyl group, a 6-quinolyl group, a 7-quinolyl group and a 8-quinolyl group) and isoquinolyl groups (a 1-isoquinolyl group, a 3-isoquinolyl group, a 4-isoquinolyl group, a 5-
15 isoquinolyl group, a 6-isoquinolyl group, a 7-isoquinolyl group and a 8-isoquinolyl group) optionally substituted with one or more of the following substituents.

Substituents: a C₁₋₆ alkyl group, a halogen atom, a C₁₋₃ alkyl group substituted with one or more fluorine
20 atoms, a nitro group, an amino group, an amino group substituted with a C₁₋₆ alkyl group, an amino group substituted with a C₁₋₆ alkylcarbonyl group, a hydroxyl group, a hydroxyl group substituted with a C₁₋₆ alkyl group, a hydroxyl group substituted with a C₁₋₆
25 alkylcarbonyl group and a C₁₋₆ alkylcarbonyl group.

Particularly preferred examples of the substituents A, R¹, R⁷ and R¹² are a phenyl group, thienyl groups (a 2-

thienyl group and a 3-thienyl group), furyl groups (a 2-furyl group and a 3-furyl group), pyridazinyl groups (a 3-pyridazinyl group and a 4-pyridazinyl group) and pyridyl groups (a 2-pyridyl group, a 3-pyridyl group and a 4-pyridyl group) optionally substituted with one or more of the following substituents.

Substituents: a C₁₋₆ alkyl group, a halogen atom, a C₁₋₃ alkyl group substituted with one or more fluorine atoms, a nitro group, an amino group, an amino group substituted with a C₁₋₆ alkyl group, an amino group substituted with an C₁₋₆ alkylcarbonyl group, a hydroxyl group, a hydroxyl group substituted with a C₁₋₆ alkyl group, a hydroxyl group substituted with a C₁₋₆ alkylcarbonyl group and a C₁₋₆ alkylcarbonyl group.

Still further preferred specific examples of the substituents A, R¹, R⁷ and R¹² are a 3-methyl-phenyl group, a 4-methyl-phenyl group, a 3,4-dimethyl-phenyl group, a 3-t-butyl-phenyl group, a 4-t-butyl-phenyl group, a 3-trifluoromethyl-phenyl group, a 4-trifluoromethyl-phenyl group, a 3,4-ditrifluoromethyl-phenyl group, a 3-chloro-phenyl group, a 4-chloro-phenyl group, a 3-iodo-phenyl group, a 4-iodo-phenyl group, a 3-fluoro-phenyl group, a 4-fluoro-phenyl group, a 3,4-dichloro-phenyl group, a 3,4-diiodo-phenyl group, a 3,4-difluoro-phenyl group, a 3-nitro-phenyl group, a 4-nitro-phenyl group, a α -naphthyl group, a β -naphthyl group and the like.

Specific preferable examples of the substituents B, R², R⁸ and R¹³ are a hydrogen atom, a methyl group, an ethyl group, a n-propyl group, an i-propyl group, a trifluoromethyl group and a phenyl group, and particularly preferred examples are a methyl group, an ethyl group and a trifluoromethyl group.

Specific preferable examples of the substituents D, R³, R⁵ and R¹⁴ are a hydrogen atom, a methyl group, an ethyl group, a n-propyl group, an i-propyl group, a c-propyl group and a phenyl group, and particularly preferable examples are a hydrogen atom, a methyl group and an ethyl group.

Specific preferable examples of the substituent R⁴ are a phenyl group, thienyl groups (a 2-thienyl group and a 3-thienyl group), furyl groups (a 2-furyl group and a 3-furyl group), pyridazinyl groups (a 3-pyridazinyl group and a 4-pyridazinyl group), pyridyl groups (a 2-pyridyl group, a 3-pyridyl group and a 4-pyridyl group), quinolyl groups (a 2-quinolyl group, a 3-quinolyl group, a 4-quinolyl group, a 5-quinolyl group, a 6-quinolyl group, a 7-quinolyl group and a 8-quinolyl group) and isoquinolyl groups (a 1-isoquinolyl group, a 3-isoquinolyl group, a 4-isoquinolyl group, a 5-isoquinolyl group, a 6-isoquinolyl group, a 7-isoquinolyl group and a 8-isoquinolyl group) substituted with one or more of the following substituents.

Substituents: a hydroxyl group, an amino group and a

nitro group.

Specific particularly preferred examples of the substituent R^4 are a phenyl group, thienyl groups (a 2-thienyl group and a 3-thienyl group), furyl groups (a 2-furyl group and a 3-furyl group), pyridazinyl groups (a 3-pyridazinyl group and a 4-pyridazinyl group) and pyridyl groups (a 2-pyridyl group, a 3-pyridyl group and a 4-pyridyl group) substituted with one or more of the following substituents.

10 Substituents: a hydroxyl group, an amino group and a nitro group.

Specific preferable example of the substituent R^{10} are a phenyl group, thienyl groups (a 2-thienyl group and a 3-thienyl group), furyl groups (a 2-furyl group and a 3-furyl group), pyridazinyl groups (a 3-pyridazinyl group and a 4-pyridazinyl group), pyridyl groups (a 2-pyridyl group, a 3-pyridyl group and a 4-pyridyl group), quinolyl groups (a 2-quinolyl group, a 3-quinolyl group, a 4-quinolyl group, a 5-quinolyl group, a 6-quinolyl group, a 7-quinolyl group and a 8-quinolyl group) and isoquinolyl groups (a 1-isoquinolyl group, a 3-isoquinolyl group, a 4-isoquinolyl group, a 5-isoquinolyl group, a 6-isoquinolyl group, a 7-isoquinolyl group and a 8-isoquinolyl group) substituted with one or more of the following substituents.

25 Substituents: a carboxyl group, sulfonic acid group, a phosphonic acid group, a carbamido group, a sulfamido

group, a hydroxycarbamido group, a hydroxysulfamido group, $\text{CH}_2\text{CO}_2\text{H}$, $\text{OCH}_2\text{CO}_2\text{H}$, $\text{NHCH}_2\text{CO}_2\text{H}$, $\text{CH}_2\text{CH}_2\text{CO}_2\text{H}$ and a tetrazole group.

Specific particularly preferred examples of the
5 substituent R^{10} are a phenyl group, thienyl groups (a 2-thienyl group and a 3-thienyl group), furyl groups (a 2-furyl group and a 3-furyl group), pyridazinyl groups (a 3-pyridazinyl group and a 4-pyridazinyl group) and
pyridyl groups (a 2-pyridyl group, a 3-pyridyl group and
10 a 4-pyridyl group) substituted with one or more of the following substituents.

Substituents: a carboxyl group, a sulfonic acid group, a phosphonic acid group, a carbamido group, a
sulfamido group, a hydroxycarbamido group, a
15 hydroxysulfamido group, $\text{CH}_2\text{CO}_2\text{H}$, $\text{OCH}_2\text{CO}_2\text{H}$, $\text{NHCH}_2\text{CO}_2\text{H}$, $\text{CH}_2\text{CH}_2\text{CO}_2\text{H}$ and a tetrazole group.

Specific preferable examples of the substituent R^{15}
are a phenyl group, thienyl groups (a 2-thienyl group and
a 3-thienyl group), furyl groups (a 2-furyl group and a
20 3-furyl group), pyridazinyl groups (a 3-pyridazinyl group
and a 4-pyridazinyl group), pyridyl groups (a 2-pyridyl
group, a 3-pyridyl group and a 4-pyridyl group), quinolyl
groups (a 2-quinolyl group, a 3-quinolyl group, a 4-
quinolyl group, a 5-quinolyl group, a 6-quinolyl group, a
25 7-quinolyl group and a 8-quinolyl group) and isoquinolyl
groups (a 1-isoquinolyl group, a 3-isoquinolyl group, a
4-isoquinolyl group, a 5-isoquinolyl group, a 6-

isoquinolyl group, a 7-isoquinolyl group and a 8-
isoquinolyl group) substituted with a substituent
selected from a hydroxyl group and an amino group and
with a substituent selected from the following
5 substituents.

Substituents: a carboxyl group, a sulfonic acid
group, a phosphonic acid group, a carbamido group, a
sulfamido group, a hydroxycarbamido group, a
hydroxysulfamido group, $\text{CH}_2\text{CO}_2\text{H}$, $\text{OCH}_2\text{CO}_2\text{H}$, $\text{NHCH}_2\text{CO}_2\text{H}$,
10 $\text{CH}_2\text{CH}_2\text{CO}_2\text{H}$ and a tetrazole group.

Specific particularly preferred examples of the
substituent R^{15} are a phenyl group, thienyl groups (a 2-
thienyl group and a 3-thienyl group), furyl groups (a 2-
furyl group and a 3-furyl group), pyridazinyl groups (a
15 3-pyridazinyl group and a 4-pyridazinyl group) and
pyridyl groups (a 2-pyridyl group, a 3-pyridyl group and
a 4-pyridyl group) substituted with a substituent
selected from a hydroxyl group and an amino group and
with a substituent selected from the following
20 substituents.

Substituents: a carboxyl group, a sulfonic acid
group, a phosphonic acid group, a carbamido group, a
sulfamido group, a hydroxycarbamido group, a
hydroxysulfamido group, $\text{CH}_2\text{CO}_2\text{H}$, $\text{OCH}_2\text{CO}_2\text{H}$, $\text{NHCH}_2\text{CO}_2\text{H}$,
25 $\text{CH}_2\text{CH}_2\text{CO}_2\text{H}$ and a tetrazole group.

Favorable compounds as the thrombopoietin receptor
activator, the preventive, therapeutic or improving agent

for diseases against which activation of the thrombopoietin receptor is effective and the platelet increasing agent of the present invention are as follows.

1) Pyrazolone compounds represented by the formula (2)

5 wherein R^4 is a C_{2-14} aryl group substituted with one or more hydroxyl groups, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

2) Pyrazolone compounds represented by the formula (2)

10 wherein R^4 is a C_{2-14} aryl group substituted with NR^5R^6 (wherein R^5 and R^6 are independently hydrogen atoms, formyl groups, C_{1-6} alkyl groups or C_{1-6} alkylcarbonyl groups), tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

15 3) Pyrazolone compounds represented by the formula (2)

wherein R^4 is a phenyl group or pyridyl group substituted with one or more hydroxyl groups, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

20 4) Pyrazolone compounds represented by the formula (2)

wherein R^4 is a phenyl group or pyridyl group substituted with NR^5R^6 (wherein R^5 and R^6 are independently hydrogen atoms, formyl groups, C_{1-6} alkyl groups or C_{1-6} alkylcarbonyl groups), tautomers, prodrugs or

25 pharmaceutically acceptable salts of the compounds or solvates thereof.

5) Pyrazolone compounds represented by the formula (2)

wherein R^4 is a thienyl group, furyl group or pyridazinyl group substituted with one or more hydroxyl groups, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

5 6) Pyrazolone compounds represented by the formula (2) wherein R^4 is a thienyl, furyl group or pyridazinyl group substituted with NR^5R^6 (wherein R^5 and R^6 are independently hydrogen atoms, formyl groups, C_{1-6} alkyl groups or C_{1-6} alkylcarbonyl groups), tautomers, prodrugs
10 or pharmaceutically acceptable salts of the compounds or solvates thereof.

7) Pyrazolone compounds represented by the formula (3) wherein R^{10} is a C_{2-14} aryl group substituted with $X(CYZ)_nCO_2H$ (wherein X is CH_2 , O, S or NR^{11} (R^{11} is a
15 hydrogen atom, a C_{1-6} alkyl group, a formyl group or a C_{1-6} alkylcarbonyl group), Y and Z are independently hydrogen atoms or C_{1-3} alkyl groups, and n is 0, 1, 2 or 3), tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

20 8) Pyrazolone compounds represented by the formula (3) wherein R^{10} is a phenyl group or pyridyl group substituted with $X(CYZ)_nCO_2H$ (wherein X is CH_2 , O, S or NR^{11} (R^{11} is a hydrogen atom, a C_{1-6} alkyl group, a formyl group or a C_{1-6} alkylcarbonyl group), Y and Z are
25 independently hydrogen atoms or C_{1-3} alkyl groups, and n is 0, 1, 2 or 3), tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

- 9) Pyrazolone compounds represented by the formula (3)
wherein R^{10} is a thienyl group, furyl group or a
pyridazinyl group substituted with $X(CYZ)_nCO_2H$ (wherein X
is CH_2 , O, S or NR^{11} (R^{11} is a hydrogen atom, a
5 C_{1-6} alkyl group, a formyl group or a C_{1-6} alkylcarbonyl
group), Y and Z are independently hydrogen atoms or C_{1-3}
alkyl groups, and n is 0, 1, 2 or 3), tautomers, prodrugs
or pharmaceutically acceptable salts of the compounds or
solvates thereof.
- 10) Pyrazolone compounds represented by the formula (3)
wherein R^{10} is a C_{2-14} aryl group substituted with a
carboxyl group, tautomers, prodrugs or pharmaceutically
acceptable salts of the compounds or solvates thereof.
- 11) Pyrazolone compounds represented by the formula (3)
15 wherein R^{10} is a phenyl group or pyridyl group
substituted with a carboxyl group, tautomers, prodrugs or
pharmaceutically acceptable salts of the compounds or
solvates thereof.
- 12) Pyrazolone compounds represented by the formula (3)
20 wherein R^{10} is a thienyl group, furyl group or
pyridazinyl group substituted with a carboxyl group,
tautomers, prodrugs or pharmaceutically acceptable salts
of the compounds or solvates thereof.
- 13) Pyrazolone compounds represented by the formula (3)
25 wherein R^{10} is a C_{2-14} aryl group substituted with a
sulfonic acid group, tautomers, prodrugs or
pharmaceutically acceptable salts of the compounds or

solvates thereof.

14) Pyrazolone compounds represented by the formula (3)
wherein R^{10} is a phenyl group or pyridyl group
substituted with a sulfonic acid group, tautomers,
5 prodrugs or pharmaceutically acceptable salts of the
compounds or solvates thereof.

15) Pyrazolone compounds represented by the formula (3)
wherein R^{10} is a thienyl group, furyl group or
pyridazinyl group substituted with a sulfonic acid group,
10 tautomers, prodrugs or pharmaceutically acceptable salts
of the compounds or solvates thereof.

16) Pyrazolone compounds represented by the formula (3)
wherein R^{10} is a C_{2-14} aryl group substituted with a
phosphonic acid group, tautomers, prodrugs or
15 pharmaceutically acceptable salts of the compounds or
solvates thereof.

17) Pyrazolone compounds represented by the formula (3)
wherein R^{10} is a phenyl group or pyridyl group
substituted with a phosphonic acid group, tautomers,
20 prodrugs or pharmaceutically acceptable salts of the
compounds or solvates thereof.

18) Pyrazolone compounds represented by the formula (3)
wherein R^{10} is a thienyl group, furyl group or
pyridazinyl group substituted with a phosphonic acid
25 group, tautomers, prodrugs or pharmaceutically acceptable
salts of the compounds or solvates thereof.

19) Pyrazolone compounds represented by the formula (3)

wherein R^{10} is a C_{2-14} aryl group substituted with a carbamido group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

20) Pyrazolone compounds represented by the formula (3)

5 wherein R^{10} is a phenyl group or pyridyl group substituted with a carbamido group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

21) Pyrazolone compounds represented by the formula (3)

10 wherein R^{10} is a thienyl group, furyl group or pyridazinyl group substituted with a carbamido group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

22) Pyrazolone compounds represented by the formula (3)

15 wherein R^{10} is a C_{2-14} aryl group substituted with a sulfamido group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

23) Pyrazolone compounds represented by the formula (3)

20 wherein R^{10} is a phenyl group or pyridyl group substituted with a sulfamido group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

24) Pyrazolone compounds represented by the formula (3)

25 wherein R^{10} is a thienyl group, furyl group or pyridazinyl group substituted with a sulfamido group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

25) Pyrazolone compounds represented by the formula (3)
wherein R^{10} is a C_{2-14} aryl group substituted with a
hydroxycarbamido group, tautomers, prodrugs or
pharmaceutically acceptable salts of the compounds or
5 solvates thereof.

26) Pyrazolone compounds represented by the formula (3)
wherein R^{10} is a phenyl group or pyridyl group
substituted with a hydroxycarbamido group, tautomers,
prodrugs or pharmaceutically acceptable salts of the
10 compounds or solvates thereof.

27) Pyrazolone compounds represented by the formula (3)
wherein R^{10} is a thienyl group, furyl group or
pyridazinyl group substituted with a hydroxycarbamido
group, tautomers, prodrugs or pharmaceutically acceptable
15 salts of the compounds or solvates thereof.

28) Pyrazolone compounds represented by the formula (3)
wherein R^{10} is a C_{2-14} aryl group substituted with a
hydroxysulfamido group, tautomers, prodrugs or
pharmaceutically acceptable salts of the compounds or
20 solvates thereof.

29) Pyrazolone compounds represented by the formula (3)
wherein R^{10} is a phenyl group or pyridyl group
substituted with a hydroxysulfamido group, tautomers,
prodrugs or pharmaceutically acceptable salts of the
25 compounds or solvates thereof.

30) Pyrazolone compounds represented by the formula (3)
wherein R^{10} is a thienyl group, furyl group or

pyridazinyl group substituted with a hydroxysulfamido group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

31) Pyrazolone compounds represented by the formula (3)
5 wherein R^{10} is a C_{2-14} aryl group substituted with a tetrazole group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

32) Pyrazolone compounds represented by the formula (3)
10 wherein R^{10} is a phenyl group or pyridyl group substituted with a tetrazole group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

33) Pyrazolone compounds represented by the formula (3)
15 wherein R^{10} is a thienyl group, furyl group or pyridazinyl group substituted with a tetrazole group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

34) Pyrazolone compounds represented by the formula (4)
20 wherein R^{15} is a C_{2-14} aryl group substituted with $X(CYZ)_nCO_2H$ (wherein X is CH_2 , O, S or NR^{16} (R^{16} is a hydrogen atom, a C_{1-6} alkyl group, a formyl group or a C_{1-6} alkylcarbonyl group), Y and Z are independently hydrogen atoms or C_{1-3} alkyl groups, and n is 0, 1, 2 or 3) and with a hydroxyl group, tautomers, prodrugs or
25 pharmaceutically acceptable salts of the compounds or solvates thereof.

35) Pyrazolone compounds represented by the formula (4)

wherein R^{15} is a phenyl or pyridyl group substituted with $X(CYZ)_nCO_2H$ (wherein X is CH_2 , O, S or NR^{16} (R^{16} is a hydrogen atom, a C_{1-6} alkyl group, a formyl group or a C_{1-6} alkylcarbonyl group), Y and Z are independently hydrogen
5 atoms or C_{1-3} alkyl groups, and n is 0, 1, 2 or 3) and with a hydroxyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

36) Pyrazolone compounds represented by the formula (4)
10 wherein R^{15} is a thienyl group, furyl group or pyridazinyl group substituted with $X(CYZ)_nCO_2H$ (wherein X is CH_2 , O, S or NR^{16} (R^{16} is a hydrogen atom, a C_{1-6} alkyl group, a formyl group or a C_{1-6} alkylcarbonyl group), Y and Z are independently hydrogen atoms or C_{1-3}
15 alkyl groups, and n is 0, 1, 2 or 3) and with a hydroxyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

37) Pyrazolone compounds represented by the formula (4)
wherein R^{15} is a C_{2-14} aryl group substituted with
20 $X(CYZ)_nCO_2H$ (wherein X is CH_2 , O, S or NR^{16} (R^{16} is a hydrogen atom, a C_{1-6} alkyl group, a formyl group or a C_{1-6} alkylcarbonyl group), Y and Z are independently hydrogen atoms or C_{1-3} alkyl groups, and n is 0, 1, 2 or 3) and with an amino group, tautomers, prodrugs or
25 pharmaceutically acceptable salts of the compounds or solvates thereof.

38) Pyrazolone compounds represented by the formula (4)

wherein R^{15} is a phenyl or pyridyl group substituted with $X(CYZ)_nCO_2H$ (wherein X is CH_2 , O, S or NR^{16} (R^{16} is a hydrogen atom, a C_{1-6} alkyl group, a formyl group or a C_{1-6} alkylcarbonyl group), Y and Z are independently hydrogen
5 atoms or C_{1-3} alkyl groups, and n is 0, 1, 2 or 3) and with an amino group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

39) Pyrazolone compounds represented by the formula (4)
10 wherein R^{15} is a thienyl group, furyl group or pyridazinyl group substituted with $X(CYZ)_nCO_2H$ (wherein X is CH_2 , O, S or NR^{16} (R^{16} is a hydrogen atom, a C_{1-6} alkyl group, a formyl group or a C_{1-6} alkylcarbonyl group), Y and Z are independently hydrogen atoms or C_{1-3}
15 alkyl groups, and n is 0, 1, 2 or 3) and with an amino group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

40) Pyrazolone compounds represented by the formula (4)
wherein R^{15} is a C_{2-14} aryl group substituted with a
20 substituent selected from a hydroxyl group, an amino group, a nitro group and a halogen atom and with a carboxyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

41) Pyrazolone compounds represented by the formula (4)
25 wherein R^{15} is a phenyl or pyridyl group substituted with a substituent selected from a hydroxyl group, an amino group, a nitro group and a halogen atom and with a

carboxyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

42) Pyrazolone compounds represented by the formula (4) wherein R^{15} is a thienyl group, furyl group or

5 pyridazinyl group substituted with a substituent selected from a hydroxyl group, an amino group, a nitro group and a halogen atom and with a carboxyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

10 43) The pyrazolone compounds according to 1), 2), 3), 4), 5) or 6), wherein R^2 is a C_{1-3} alkyl group substituted with one or more fluorine atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

15 44) The pyrazolone compounds according to 1), 2), 3), 4), 5) or 6), wherein R^2 is a C_{1-6} alkyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

45) The pyrazolone compounds according to 1), 2), 3), 4),
20 5) or 6), wherein R^2 is hydrogen, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

46) The pyrazolone compounds according to 7), 8), 9), 10), 11), 12), 13) 14), 15), 16), 17), 18), 19), 20), 21), 22),
25 23), 24), 25), 26), 27), 28), 29), 30), 31), 32) or 33), wherein R^8 is a C_{1-3} alkyl group substituted with one or more fluorine atoms, tautomers, prodrugs or

pharmaceutically acceptable salts of the compounds or solvates thereof.

47) The pyrazolone compounds according to 7), 8), 9), 10), 11), 12), 13) 14), 15), 16), 17), 18), 19), 20), 21), 22), 5 23), 24), 25), 26), 27), 28), 29), 30), 31), 32) or 33), wherein R^8 is a C_{1-6} alkyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

48) The pyrazolone compounds according to 7), 8), 9), 10), 10 11), 12), 13) 14), 15), 16), 17), 18), 19), 20), 21), 22), 23), 24), 25), 26), 27), 28), 29), 30), 31), 32) or 33), wherein R^8 is hydrogen, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

15 49) The pyrazolone compounds according to 34), 35), 36), 37), 38), 39), 40), 41) or 42), wherein R^{13} is a C_{1-3} alkyl group substituted with one or more fluorine atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

20 50) The pyrazolone compounds according to 34), 35), 36), 37), 38), 39), 40), 41) or 42), wherein R^{13} is a C_{1-6} alkyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

51) The pyrazolone compounds according to 34), 35), 36), 25 37), 38), 39), 40), 41) or 42), wherein R^{13} is hydrogen, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

52) The pyrazolone compounds according to 1), 2), 3), 4), 5), 6), 43), 44) or 45), wherein R^3 is a hydrogen atom, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

5 53) The pyrazolone compounds according to 1), 2), 3), 4), 5), 6), 43), 44) or 45), wherein R^3 is a C_{1-6} alkyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

54) The pyrazolone compounds according to 7), 8), 9), 10),
10 11), 12), 13), 14), 15), 16), 17), 18), 19), 20), 21), 22), 23), 24), 25), 26), 27), 28), 29), 30), 31), 32), 33), 46), 47) or 48), wherein R^9 is a hydrogen atom, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

15 55) The pyrazolone compounds according to 7), 8), 9), 10), 11), 12), 13), 14), 15), 16), 17), 18), 19), 20), 21), 22), 23), 24), 25), 26), 27), 28), 29), 30), 31), 32), 33), 46), 47) or 48), wherein R^9 is a C_{1-6} alkyl group, tautomers, prodrugs or pharmaceutically acceptable salts
20 of the compounds or solvates thereof.

56) The pyrazolone compounds according to 34), 35), 36), 37), 38), 39), 40), 41), 42), 49), 50) or 51), wherein R^{14} is a hydrogen atom, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or
25 solvates thereof.

57) The pyrazolone compounds according to 34), 35), 36), 37), 38), 39), 40), 41), 42), 49), 50) or 51), wherein

R¹⁴ is a C₁₋₆ alkyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

58) The pyrazolone compounds according to 52) or 53),
5 wherein R¹ is a C₂₋₁₄ aryl group substituted with one or more C₁₋₆ alkyl groups, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

59) The pyrazolone compounds according to 52) or 53),
10 wherein R¹ is a phenyl group or pyridyl group substituted with one or more C₁₋₆ alkyl groups, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

60) The pyrazolone compounds according to 52) or 53),
15 wherein R¹ is a thienyl group, furyl group or pyridazinyl group substituted with one or more C₁₋₆ alkyl groups, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

61) The pyrazolone compounds according to 54) or 55),
20 wherein R⁷ is a C₂₋₁₄ aryl group substituted with one or more C₁₋₆ alkyl groups, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

62) The pyrazolone compounds according to 54) or 55),
25 wherein R⁷ is a phenyl group or pyridyl group substituted with one or more C₁₋₆ alkyl groups, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or

solvates thereof.

63) The pyrazolone compounds according to 54) or 55),
wherein R^7 is a thienyl group, furyl group or pyridazinyl
group substituted with one or more C_{1-6} alkyl groups,
5 tautomers, prodrugs or pharmaceutically acceptable salts
of the compounds or solvates thereof.

64) The pyrazolone compounds according to 56) or 57),
wherein R^{12} is a C_{2-14} aryl group substituted with one or
more C_{1-6} alkyl groups, tautomers, prodrugs or
10 pharmaceutically acceptable salts of the compounds or
solvates thereof.

65) The pyrazolone compounds according to 56) or 57),
wherein R^{12} is a phenyl group or pyridyl group
substituted with one or more C_{1-6} alkyl groups, tautomers,
15 prodrugs or pharmaceutically acceptable salts of the
compounds or solvates thereof.

66) The pyrazolone compounds according to 56) or 57),
wherein R^{12} is a thienyl group, furyl group or
pyridazinyl group substituted with one or more C_{1-6} alkyl
20 groups, tautomers, prodrugs or pharmaceutically
acceptable salts of the compounds or solvates thereof.

67) The pyrazolone compounds according to 52) or 53),
wherein R^1 is a C_{2-14} aryl group substituted with one or
more halogen atoms, tautomers, prodrugs or
25 pharmaceutically acceptable salts of the compounds or
solvates thereof.

68) The pyrazolone compounds according to 52) or 53),

wherein R^1 is a phenyl group or pyridyl group substituted with one or more halogen atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

5 69) The pyrazolone compounds according to 52) or 53), wherein R^1 is a thienyl group, furyl group or pyridazinyl group substituted with one or more halogen atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

10 70) The pyrazolone compounds according to 54) or 55), wherein R^7 is a C_{2-14} aryl group substituted with one or more halogen atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

15 71) The pyrazolone compounds according to 54) or 55), wherein R^7 is a phenyl group or pyridyl group substituted with one or more halogen atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

20 72) The pyrazolone compounds according to 54) or 55), wherein R^7 is a thienyl group, furyl group or pyridazinyl group substituted with one or more halogen atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

25 73) The pyrazolone compounds according to 56) or 57), wherein R^{12} is a C_{2-14} aryl group substituted with one or more halogen atoms, tautomers, prodrugs or

pharmaceutically acceptable salts of the compounds or solvates thereof.

74) The pyrazolone compounds according to 56) or 57), wherein R^{12} is a phenyl group or pyridyl group substituted with one or more halogen atoms, tautomers, 5 prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

75) The pyrazolone compounds according to 56) or 57), wherein R^{12} is a thienyl group, furyl group or 10 pyridazinyl group substituted with one or more halogen atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

76) The pyrazolone compounds according to 52) or 53), wherein R^1 is a C_{2-14} aryl group substituted with one or 15 more C_{1-3} alkyl groups substituted with one or more fluorine atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

77) The pyrazolone compounds according to 52) or 53), wherein R^1 is a phenyl group or pyridyl group substituted 20 with one or more C_{1-3} alkyl groups substituted with one or more fluorine atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

78) The pyrazolone compounds according to 52) or 53), 25 wherein R^1 is a thienyl group, furyl group or pyridazinyl group substituted with one or more C_{1-3} alkyl groups substituted with one or more fluorine atoms, tautomers,

prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

79) The pyrazolone compounds according to 54) or 55), wherein R⁷ is a C₂₋₁₄ aryl group substituted with one or
5 more C₁₋₃ alkyl groups substituted with one or more fluorine atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

80) The pyrazolone compounds according to 54) or 55), wherein R⁷ is a phenyl group or pyridyl group substituted
10 with one or more C₁₋₃ alkyl groups substituted with one or more fluorine atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

81) The pyrazolone compounds according to 54) or 55),
15 wherein R⁷ is a thienyl group, furyl group or pyridazinyl group substituted with one or more C₁₋₃ alkyl groups substituted with one or more fluorine atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

20 82) The pyrazolone compounds according to 56) or 57), wherein R¹² is a C₂₋₁₄ aryl group substituted with one or more C₁₋₃ alkyl groups substituted with one or more fluorine atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

25 83) The pyrazolone compounds according to 56) or 57), wherein R¹² is a phenyl group or pyridyl group substituted with one or more C₁₋₃ alkyl groups substituted

with one or more fluorine atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

84) The pyrazolone compounds according to 56) or 57),
5 wherein R^{12} is a thienyl group, furyl group or pyridazinyl group substituted with one or more C_{1-3} alkyl groups substituted with one or more fluorine atoms, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

10 85) The pyrazolone compounds according to 52) or 53), wherein R^1 is a C_{2-14} aryl group substituted with a hydroxyl group substituted with a C_{1-6} alkyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

15 86) The pyrazolone compounds according to 52) or 53), wherein R^1 is a phenyl group or pyridyl group substituted with a hydroxyl group substituted with a C_{1-6} alkyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

20 87) The pyrazolone compounds according to 52) or 53), wherein R^1 is a thienyl group, furyl group or pyridazinyl group substituted with a hydroxyl group substituted with a C_{1-6} alkyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or
25 solvates thereof.

88) The pyrazolone compounds according to 54) or 55), wherein R^7 is a C_{2-14} aryl group substituted with a

hydroxyl group substituted with a C₁₋₆ alkyl group,
tautomers, prodrugs or pharmaceutically acceptable salts
of the compounds or solvates thereof.

89) The pyrazolone compounds according to 54) or 55),
5 wherein R⁷ is a phenyl group or pyridyl group substituted
with a hydroxyl group substituted with a C₁₋₆ alkyl group,
tautomers, prodrugs or pharmaceutically acceptable salts
of the compounds or solvates thereof.

90) The pyrazolone compounds according to 54) or 55),
10 wherein R⁷ is a thienyl group, furyl group or pyridazinyl
group substituted with a hydroxyl group substituted with
a C₁₋₆ alkyl group, tautomers, prodrugs or
pharmaceutically acceptable salts of the compounds or
solvates thereof.

15 91) The pyrazolone compounds according to 56) or 57),
wherein R¹² is a C₂₋₁₄ aryl group substituted with a
hydroxyl group substituted with a C₁₋₆ alkyl group,
tautomers, prodrugs or pharmaceutically acceptable salts
of the compounds or solvates thereof.

20 92) The pyrazolone compounds according to 56) or 57),
wherein R¹² is a phenyl group or pyridyl group
substituted with a hydroxyl group substituted with a C₁₋₆
alkyl group, tautomers, prodrugs or pharmaceutically
acceptable salts of the compounds or solvates thereof.

25 93) The pyrazolone compounds according to 56) or 57),
wherein R¹² is a thienyl group, furyl group or
pyridazinyl group substituted with a hydroxyl group

substituted with a C₁₋₆ alkyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

94) The pyrazolone compounds according to 52) or 53),
5 wherein R¹ is a C₂₋₁₄ aryl group substituted with an amino group substituted with a C₁₋₆ alkyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

95) The pyrazolone compounds according to 52) or 53),
10 wherein R¹ is a phenyl group or pyridyl group substituted with an amino group substituted with a C₁₋₆ alkyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

96) The pyrazolone compounds according to 52) or 53),
15 wherein R¹ is a thienyl group, furyl group or pyridazinyl group substituted with an amino group substituted with a C₁₋₆ alkyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

97) The pyrazolone compounds according to 54) or 55),
20 wherein R⁷ is a C₂₋₁₄ aryl group substituted with an amino group substituted with a C₁₋₆ alkyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

98) The pyrazolone compounds according to 54) or 55),
25 wherein R⁷ is a phenyl group or pyridyl group substituted with an amino group substituted with a C₁₋₆ alkyl group, tautomers, prodrugs or pharmaceutically acceptable salts

of the compounds or solvates thereof.

99) The pyrazolone compounds according to 54) or 55),
wherein R⁷ is a thienyl group, furyl group or pyridazinyl
group substituted with an amino group substituted with a
5 C₁₋₆ alkyl group, tautomers, prodrugs or pharmaceutically
acceptable salts of the compounds or solvates thereof.

100) The pyrazolone compounds according to 56) or 57),
wherein R¹² is a C₂₋₁₄ aryl group substituted with an amino
group substituted with a C₁₋₆ alkyl group, tautomers,
10 prodrugs or pharmaceutically acceptable salts of the
compounds or solvates thereof.

101) The pyrazolone compounds according to 56) or 57),
wherein R¹² is a phenyl group or pyridyl group
substituted with an amino group substituted with a C₁₋₆
15 alkyl group, tautomers, prodrugs or pharmaceutically
acceptable salts of the compounds or solvates thereof.

102) The pyrazolone compounds according to 56) or 57),
wherein R¹² is a thienyl group, furyl group or
pyridazinyl group substituted with an amino group
20 substituted with a C₁₋₆ alkyl group, tautomers, prodrugs
or pharmaceutically acceptable salts of the compounds or
solvates thereof.

103) The pyrazolone compounds according to 52) or 53),
wherein R¹ is a C₂₋₁₄ aryl group substituted with a
25 hydroxyl group substituted with a C₁₋₆ alkylcarbonyl group,
tautomers, prodrugs or pharmaceutically acceptable salts
of the compounds or solvates thereof.

- 104) The pyrazolone compounds according to 52) or 53),
wherein R¹ is a phenyl group or pyridyl group substituted
with a hydroxyl group substituted with a C₁₋₆
alkylcarbonyl group, tautomers, prodrugs or
5 pharmaceutically acceptable salts of the compounds or
solvates thereof.
- 105) The pyrazolone compounds according to 52) or 53),
wherein R¹ is a thienyl group, furyl group or pyridazinyl
group substituted with a hydroxyl group substituted with
10 a C₁₋₆ alkylcarbonyl group, tautomers, prodrugs or
pharmaceutically acceptable salts of the compounds or
solvates thereof.
- 106) The pyrazolone compounds according to 54) or 55),
wherein R⁷ is a C₂₋₁₄ aryl group substituted with a
15 hydroxyl group substituted with a C₁₋₆ alkylcarbonyl group,
tautomers, prodrugs or pharmaceutically acceptable salts
of the compounds or solvates thereof.
- 107) The pyrazolone compounds according to 54) or 55),
wherein R⁷ is a phenyl group or pyridyl group substituted
20 with a hydroxyl group substituted with a C₁₋₆
alkylcarbonyl group, tautomers, prodrugs or
pharmaceutically acceptable salts of the compounds or
solvates thereof.
- 108) The pyrazolone compounds according to 54) or 55),
25 wherein R⁷ is a thienyl group, furyl group or pyridazinyl
group substituted with a hydroxyl group substituted with
a C₁₋₆ alkylcarbonyl group, tautomers, prodrugs or

pharmaceutically acceptable salts of the compounds or solvates thereof.

109) The pyrazolone compounds according to 56) or 57), wherein R^{12} is a C_{2-14} aryl group substituted with a hydroxyl group substituted with a C_{1-6} alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

110) The pyrazolone compounds according to 56) or 57), wherein R^{12} is a phenyl group or pyridyl group substituted with a hydroxyl group substituted with a C_{1-6} alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

111) The pyrazolone compounds according to 56) or 57), wherein R^{12} is a thienyl group, furyl group or pyridazinyl group substituted with a hydroxyl group substituted with a C_{1-6} alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

112) The pyrazolone compounds according to 52) or 53), wherein R^1 is a C_{2-14} aryl group substituted with an amino group substituted with a C_{1-6} alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

113) The pyrazolone compounds according to 52) or 53), wherein R^1 is a phenyl group or pyridyl group substituted with an amino group substituted with a C_{1-6} alkylcarbonyl

group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

114) The pyrazolone compounds according to 52) or 53), wherein R^1 is a thienyl group, furyl group or pyridazinyl group substituted with an amino group substituted with a C_{1-6} alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

115) The pyrazolone compounds according to 54) or 55), wherein R^7 is a C_{2-14} aryl group substituted with an amino group substituted with a C_{1-6} alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

116) The pyrazolone compounds according to 54) or 55), wherein R^7 is a phenyl group or pyridyl group substituted with an amino group substituted with a C_{1-6} alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

117) The pyrazolone compounds according to 54) or 55), wherein R^7 is a thienyl group, furyl group or pyridazinyl group substituted with an amino group substituted with a C_{1-6} alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

118) The pyrazolone compounds according to 56) or 57), wherein R^{12} is a C_{2-14} aryl group substituted with an amino group substituted with a C_{1-6} alkylcarbonyl group,

tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

119) The pyrazolone compounds according to 56) or 57), wherein R^{12} is a phenyl group or pyridyl group substituted with an amino group substituted with a C_{1-6} alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

120) The pyrazolone compounds according to 56) or 57), wherein R^{12} is a thienyl group, furyl group or pyridazinyl group substituted with an amino group substituted with a C_{1-6} alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

121) The pyrazolone compounds according to 52) or 53), wherein R^1 is a C_{2-14} aryl group substituted with a C_{1-6} alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

122) The pyrazolone compounds according to 52) or 53), wherein R^1 is a phenyl group or pyridyl group substituted with a C_{1-6} alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

123) The pyrazolone compounds according to 52) or 53), wherein R^1 is a thienyl group, furyl group or pyridazinyl group substituted with a C_{1-6} alkylcarbonyl group,

tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

124) The pyrazolone compounds according to 54) or 55), wherein R⁷ is a C₂₋₁₄ aryl group substituted with a C₁₋₆ alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

125) The pyrazolone compounds according to 54) or 55), wherein R⁷ is a phenyl group or pyridyl group substituted with a C₁₋₆ alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

126) The pyrazolone compounds according to 54) or 55), wherein R⁷ is a thienyl group, furyl group or pyridazinyl group substituted with a C₁₋₆ alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

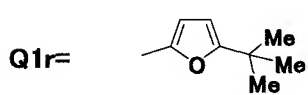
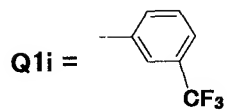
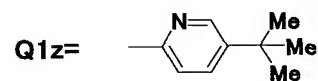
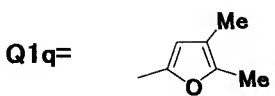
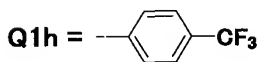
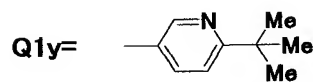
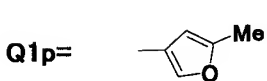
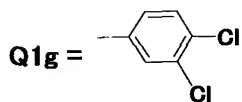
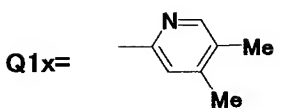
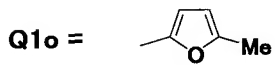
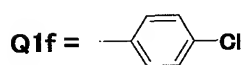
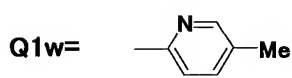
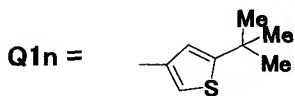
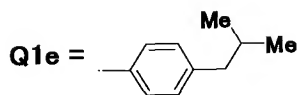
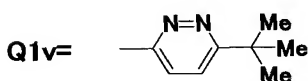
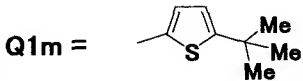
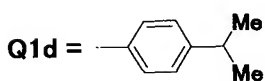
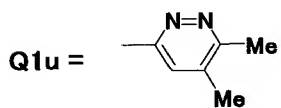
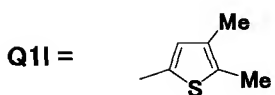
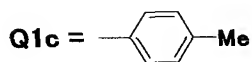
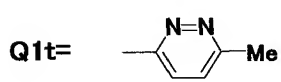
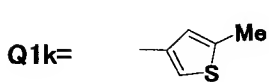
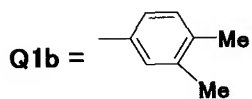
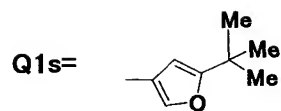
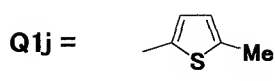
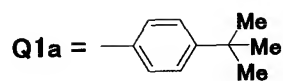
127) The pyrazolone compounds according to 56) or 57), wherein R¹² is a C₂₋₁₄ aryl group substituted with a C₁₋₆ alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

128) The pyrazolone compounds according to 56) or 57), wherein R¹² is a phenyl group or pyridyl group substituted with a C₁₋₆ alkylcarbonyl group, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof.

129) The pyrazolone compounds according to 56) or 57),
wherein R¹² is a thienyl group, furyl group or
pyridazinyl group substituted with a C₁₋₆ alkylcarbonyl
group, tautomers, prodrugs or pharmaceutically acceptable
5 salts of the compounds or solvates thereof.

130) The compounds wherein R⁷, R⁸, R⁹ and R¹⁰ are any of
the following combinations in Table 1, tautomers,
prodrugs or pharmaceutically acceptable salts of the
compounds or solvates thereof. The symbols in Table 1
10 denote the following substituents.

【Ka 9】



【Ka 10】

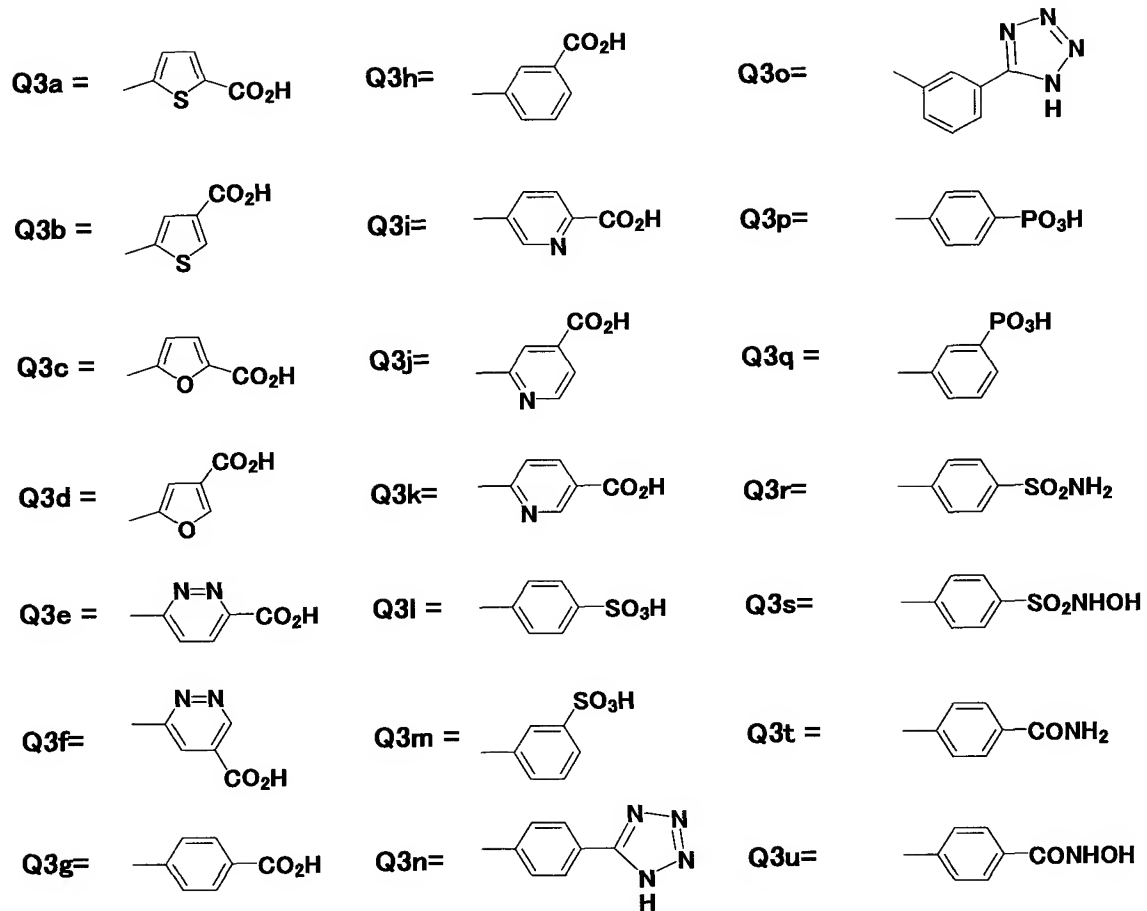


Table 1

No	R ⁷	R ⁸	R ⁹	R ¹⁰
1	Q1a	H	H	Q3a
2	Q1a	H	H	Q3b
3	Q1a	H	H	Q3c
4	Q1a	H	H	Q3d
5	Q1a	H	H	Q3e
6	Q1a	H	H	Q3f
7	Q1a	H	H	Q3g
8	Q1a	H	H	Q3h
9	Q1a	H	H	Q3i
10	Q1a	H	H	Q3j
11	Q1a	H	H	Q3k
12	Q1a	H	H	Q3l
13	Q1a	H	H	Q3m
14	Q1a	H	H	Q3n
15	Q1a	H	H	Q3o
16	Q1a	H	H	Q3p
17	Q1a	H	H	Q3q
18	Q1a	H	Me	Q3a
19	Q1a	H	Me	Q3b
20	Q1a	H	Me	Q3c
21	Q1a	H	Me	Q3d
22	Q1a	H	Me	Q3e
23	Q1a	H	Me	Q3f
24	Q1a	H	Me	Q3g
25	Q1a	H	Me	Q3h
26	Q1a	H	Me	Q3i
27	Q1a	H	Me	Q3j
28	Q1a	H	Me	Q3k
29	Q1a	H	Me	Q3l
30	Q1a	H	Me	Q3m
31	Q1a	H	Me	Q3n
32	Q1a	H	Me	Q3o
33	Q1a	H	Me	Q3p

34	Q1a	H	Me	Q3q
35	Q1a	Me	H	Q3a
36	Q1a	Me	H	Q3b
37	Q1a	Me	H	Q3c
38	Q1a	Me	H	Q3d
39	Q1a	Me	H	Q3e
40	Q1a	Me	H	Q3f
41	Q1a	Me	H	Q3g
42	Q1a	Me	H	Q3h
43	Q1a	Me	H	Q3i
44	Q1a	Me	H	Q3j
45	Q1a	Me	H	Q3k
46	Q1a	Me	H	Q3l
47	Q1a	Me	H	Q3m
48	Q1a	Me	H	Q3n
49	Q1a	Me	H	Q3o
50	Q1a	Me	H	Q3p
51	Q1a	Me	H	Q3q
52	Q1a	Me	Me	Q3a
53	Q1a	Me	Me	Q3b
54	Q1a	Me	Me	Q3c
55	Q1a	Me	Me	Q3d
56	Q1a	Me	Me	Q3e
57	Q1a	Me	Me	Q3f
58	Q1a	Me	Me	Q3g
59	Q1a	Me	Me	Q3h
60	Q1a	Me	Me	Q3i
61	Q1a	Me	Me	Q3j
62	Q1a	Me	Me	Q3k
63	Q1a	Me	Me	Q3l
64	Q1a	Me	Me	Q3m
65	Q1a	Me	Me	Q3n
66	Q1a	Me	Me	Q3o
67	Q1a	Me	Me	Q3p
68	Q1a	Me	Me	Q3q
69	Q1a	CF3	H	Q3a

70	Q1a	CF3	H	Q3b
71	Q1a	CF3	H	Q3c
72	Q1a	CF3	H	Q3d
73	Q1a	CF3	H	Q3e
74	Q1a	CF3	H	Q3f
75	Q1a	CF3	H	Q3g
76	Q1a	CF3	H	Q3h
77	Q1a	CF3	H	Q3i
78	Q1a	CF3	H	Q3j
79	Q1a	CF3	H	Q3k
80	Q1a	CF3	H	Q3l
81	Q1a	CF3	H	Q3m
82	Q1a	CF3	H	Q3n
83	Q1a	CF3	H	Q3o
84	Q1a	CF3	H	Q3p
85	Q1a	CF3	H	Q3q
86	Q1a	CF3	Me	Q3a
87	Q1a	CF3	Me	Q3b
88	Q1a	CF3	Me	Q3c
89	Q1a	CF3	Me	Q3d
90	Q1a	CF3	Me	Q3e
91	Q1a	CF3	Me	Q3f
92	Q1a	CF3	Me	Q3g
93	Q1a	CF3	Me	Q3h
94	Q1a	CF3	Me	Q3i
95	Q1a	CF3	Me	Q3j
96	Q1a	CF3	Me	Q3k
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101	Q1a	CF3	Me	Q3p
102	Q1a	CF3	Me	Q3q
103	Q1b	H	H	Q3a
104	Q1b	H	H	Q3b
105	Q1b	H	H	Q3c

106	Q1b	H	H	Q3d
107	Q1b	H	H	Q3e
108	Q1b	H	H	Q3f
109	Q1b	H	H	Q3g
110	Q1b	H	H	Q3h
111	Q1b	H	H	Q3i
112	Q1b	H	H	Q3j
113	Q1b	H	H	Q3k
114	Q1b	H	H	Q3l
115	Q1b	H	H	Q3m
116	Q1b	H	H	Q3n
117	Q1b	H	H	Q3o
118	Q1b	H	H	Q3p
119	Q1b	H	H	Q3q
120	Q1b	H	Me	Q3a
121	Q1b	H	Me	Q3b
122	Q1b	H	Me	Q3c
123	Q1b	H	Me	Q3d
124	Q1b	H	Me	Q3e
125	Q1b	H	Me	Q3f
126	Q1b	H	Me	Q3g
127	Q1b	H	Me	Q3h
128	Q1b	H	Me	Q3i
129	Q1b	H	Me	Q3j
130	Q1b	H	Me	Q3k
131	Q1b	H	Me	Q3l
132	Q1b	H	Me	Q3m
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136	Q1b	H	Me	Q3q
137	Q1b	Me	H	Q3a
138	Q1b	Me	H	Q3b
139	Q1b	Me	H	Q3c
140	Q1b	Me	H	Q3d
141	Q1b	Me	H	Q3e

142	Q1b	Me	H	Q3f
143	Q1b	Me	H	Q3g
144	Q1b	Me	H	Q3h
145	Q1b	Me	H	Q3i
146	Q1b	Me	H	Q3j
147	Q1b	Me	H	Q3k
148	Q1b	Me	H	Q3l
149	Q1b	Me	H	Q3m
150	Q1b	Me	H	Q3n
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152	Q1b	Me	H	Q3p
153	Q1b	Me	H	Q3q
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155	Q1b	Me	Me	Q3b
156	Q1b	Me	Me	Q3c
157	Q1b	Me	Me	Q3d
158	Q1b	Me	Me	Q3e
159	Q1b	Me	Me	Q3f
160	Q1b	Me	Me	Q3g
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194	Q1b	CF3	Me	Q3g
195	Q1b	CF3	Me	Q3h
196	Q1b	CF3	Me	Q3i
197	Q1b	CF3	Me	Q3j
198	Q1b	CF3	Me	Q3k
199	Q1b	CF3	Me	Q3l
200	Q1b	CF3	Me	Q3m
201	Q1b	CF3	Me	Q3n
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203	Q1b	CF3	Me	Q3p
204	Q1b	CF3	Me	Q3q
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209	Q1c	H	H	Q3e
210	Q1c	H	H	Q3f
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245	Q1c	Me	H	Q3g
246	Q1c	Me	H	Q3h
247	Q1c	Me	H	Q3i
248	Q1c	Me	H	Q3j
249	Q1c	Me	H	Q3k

250	Q1c	Me	H	Q3l
251	Q1c	Me	H	Q3m
252	Q1c	Me	H	Q3n
253	Q1c	Me	H	Q3o
254	Q1c	Me	H	Q3p
255	Q1c	Me	H	Q3q
256	Q1c	Me	Me	Q3a
257	Q1c	Me	Me	Q3b
258	Q1c	Me	Me	Q3c
259	Q1c	Me	Me	Q3d
260	Q1c	Me	Me	Q3e
261	Q1c	Me	Me	Q3f
262	Q1c	Me	Me	Q3g
263	Q1c	Me	Me	Q3h
264	Q1c	Me	Me	Q3i
265	Q1c	Me	Me	Q3j
266	Q1c	Me	Me	Q3k
267	Q1c	Me	Me	Q3l
268	Q1c	Me	Me	Q3m
269	Q1c	Me	Me	Q3n
270	Q1c	Me	Me	Q3o
271	Q1c	Me	Me	Q3p
272	Q1c	Me	Me	Q3q
273	Q1c	CF3	H	Q3a
274	Q1c	CF3	H	Q3b
275	Q1c	CF3	H	Q3c
276	Q1c	CF3	H	Q3d
277	Q1c	CF3	H	Q3e
278	Q1c	CF3	H	Q3f
279	Q1c	CF3	H	Q3g
280	Q1c	CF3	H	Q3h
281	Q1c	CF3	H	Q3i
282	Q1c	CF3	H	Q3j
283	Q1c	CF3	H	Q3k
284	Q1c	CF3	H	Q3l
285	Q1c	CF3	H	Q3m

286	Q1c	CF3	H	Q3n
287	Q1c	CF3	H	Q3o
288	Q1c	CF3	H	Q3p
289	Q1c	CF3	H	Q3q
290	Q1c	CF3	Me	Q3a
291	Q1c	CF3	Me	Q3b
292	Q1c	CF3	Me	Q3c
293	Q1c	CF3	Me	Q3d
294	Q1c	CF3	Me	Q3e
295	Q1c	CF3	Me	Q3f
296	Q1c	CF3	Me	Q3g
297	Q1c	CF3	Me	Q3h
298	Q1c	CF3	Me	Q3i
299	Q1c	CF3	Me	Q3j
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301	Q1c	CF3	Me	Q3l
302	Q1c	CF3	Me	Q3m
303	Q1c	CF3	Me	Q3n
304	Q1c	CF3	Me	Q3o
305	Q1c	CF3	Me	Q3p
306	Q1c	CF3	Me	Q3q
307	Q1d	H	H	Q3a
308	Q1d	H	H	Q3b
309	Q1d	H	H	Q3c
310	Q1d	H	H	Q3d
311	Q1d	H	H	Q3e
312	Q1d	H	H	Q3f
313	Q1d	H	H	Q3g
314	Q1d	H	H	Q3h
315	Q1d	H	H	Q3i
316	Q1d	H	H	Q3j
317	Q1d	H	H	Q3k
318	Q1d	H	H	Q3l
319	Q1d	H	H	Q3m
320	Q1d	H	H	Q3n
321	Q1d	H	H	Q3o

322	Q1d	H	H	Q3p
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324	Q1d	H	Me	Q3a
325	Q1d	H	Me	Q3b
326	Q1d	H	Me	Q3c
327	Q1d	H	Me	Q3d
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331	Q1d	H	Me	Q3h
332	Q1d	H	Me	Q3i
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334	Q1d	H	Me	Q3k
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336	Q1d	H	Me	Q3m
337	Q1d	H	Me	Q3n
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339	Q1d	H	Me	Q3p
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342	Q1d	Me	H	Q3b
343	Q1d	Me	H	Q3c
344	Q1d	Me	H	Q3d
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347	Q1d	Me	H	Q3g
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355	Q1d	Me	H	Q3o
356	Q1d	Me	H	Q3p
357	Q1d	Me	H	Q3q

358	Q1d	Me	Me	Q3a
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361	Q1d	Me	Me	Q3d
362	Q1d	Me	Me	Q3e
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375	Q1d	CF3	H	Q3a
376	Q1d	CF3	H	Q3b
377	Q1d	CF3	H	Q3c
378	Q1d	CF3	H	Q3d
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381	Q1d	CF3	H	Q3g
382	Q1d	CF3	H	Q3h
383	Q1d	CF3	H	Q3i
384	Q1d	CF3	H	Q3j
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386	Q1d	CF3	H	Q3l
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389	Q1d	CF3	H	Q3o
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391	Q1d	CF3	H	Q3q
392	Q1d	CF3	Me	Q3a
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394	Q1d	CF3	Me	Q3c
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396	Q1d	CF3	Me	Q3e
397	Q1d	CF3	Me	Q3f
398	Q1d	CF3	Me	Q3g
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403	Q1d	CF3	Me	Q3l
404	Q1d	CF3	Me	Q3m
405	Q1d	CF3	Me	Q3n
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408	Q1d	CF3	Me	Q3q
409	Q1e	H	H	Q3a
410	Q1e	H	H	Q3b
411	Q1e	H	H	Q3c
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413	Q1e	H	H	Q3e
414	Q1e	H	H	Q3f
415	Q1e	H	H	Q3g
416	Q1e	H	H	Q3h
417	Q1e	H	H	Q3i
418	Q1e	H	H	Q3j
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420	Q1e	H	H	Q3l
421	Q1e	H	H	Q3m
422	Q1e	H	H	Q3n
423	Q1e	H	H	Q3o
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425	Q1e	H	H	Q3q
426	Q1e	H	Me	Q3a
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476	Q1e	Me	Me	Q3q
477	Q1e	CF3	H	Q3a
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495	Q1e	CF3	Me	Q3b
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554	Q1f	Me	H	Q3j
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561	Q1f	Me	H	Q3q
562	Q1f	Me	Me	Q3a
563	Q1f	Me	Me	Q3b
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566	Q1f	Me	Me	Q3e
567	Q1f	Me	Me	Q3f
568	Q1f	Me	Me	Q3g
569	Q1f	Me	Me	Q3h
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571	Q1f	Me	Me	Q3j
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573	Q1f	Me	Me	Q3l

574	Q1f	Me	Me	Q3m
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576	Q1f	Me	Me	Q3o
577	Q1f	Me	Me	Q3p
578	Q1f	Me	Me	Q3q
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580	Q1f	CF3	H	Q3b
581	Q1f	CF3	H	Q3c
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612	Q1f	CF3	Me	Q3q
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614	Q1g	H	H	Q3b
615	Q1g	H	H	Q3c
616	Q1g	H	H	Q3d
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618	Q1g	H	H	Q3f
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624	Q1g	H	H	Q3l
625	Q1g	H	H	Q3m
626	Q1g	H	H	Q3n
627	Q1g	H	H	Q3o
628	Q1g	H	H	Q3p
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630	Q1g	H	Me	Q3a
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632	Q1g	H	Me	Q3c
633	Q1g	H	Me	Q3d
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636	Q1g	H	Me	Q3g
637	Q1g	H	Me	Q3h
638	Q1g	H	Me	Q3i
639	Q1g	H	Me	Q3j
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641	Q1g	H	Me	Q3l
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643	Q1g	H	Me	Q3n
644	Q1g	H	Me	Q3o
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649	Q1g	Me	H	Q3c
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672	Q1g	Me	Me	Q3i
673	Q1g	Me	Me	Q3j
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677	Q1g	Me	Me	Q3n
678	Q1g	Me	Me	Q3o
679	Q1g	Me	Me	Q3p
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681	Q1g	CF3	H	Q3a

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684	Q1g	CF3	H	Q3d
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699	Q1g	CF3	Me	Q3b
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709	Q1g	CF3	Me	Q3l
710	Q1g	CF3	Me	Q3m
711	Q1g	CF3	Me	Q3n
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713	Q1g	CF3	Me	Q3p
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715	Q1h	H	H	Q3a
716	Q1h	H	H	Q3b
717	Q1h	H	H	Q3c

718	Q1h	H	H	Q3d
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721	Q1h	H	H	Q3g
722	Q1h	H	H	Q3h
723	Q1h	H	H	Q3i
724	Q1h	H	H	Q3j
725	Q1h	H	H	Q3k
726	Q1h	H	H	Q3l
727	Q1h	H	H	Q3m
728	Q1h	H	H	Q3n
729	Q1h	H	H	Q3o
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731	Q1h	H	H	Q3q
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734	Q1h	H	Me	Q3c
735	Q1h	H	Me	Q3d
736	Q1h	H	Me	Q3e
737	Q1h	H	Me	Q3f
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747	Q1h	H	Me	Q3p
748	Q1h	H	Me	Q3q
749	Q1h	Me	H	Q3a
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751	Q1h	Me	H	Q3c
752	Q1h	Me	H	Q3d
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757	Q1h	Me	H	Q3i
758	Q1h	Me	H	Q3j
759	Q1h	Me	H	Q3k
760	Q1h	Me	H	Q3l
761	Q1h	Me	H	Q3m
762	Q1h	Me	H	Q3n
763	Q1h	Me	H	Q3o
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765	Q1h	Me	H	Q3q
766	Q1h	Me	Me	Q3a
767	Q1h	Me	Me	Q3b
768	Q1h	Me	Me	Q3c
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770	Q1h	Me	Me	Q3e
771	Q1h	Me	Me	Q3f
772	Q1h	Me	Me	Q3g
773	Q1h	Me	Me	Q3h
774	Q1h	Me	Me	Q3i
775	Q1h	Me	Me	Q3j
776	Q1h	Me	Me	Q3k
777	Q1h	Me	Me	Q3l
778	Q1h	Me	Me	Q3m
779	Q1h	Me	Me	Q3n
780	Q1h	Me	Me	Q3o
781	Q1h	Me	Me	Q3p
782	Q1h	Me	Me	Q3q
783	Q1h	CF3	H	Q3a
784	Q1h	CF3	H	Q3b
785	Q1h	CF3	H	Q3c
786	Q1h	CF3	H	Q3d
787	Q1h	CF3	H	Q3e
788	Q1h	CF3	H	Q3f
789	Q1h	CF3	H	Q3g

790	Q1h	CF3	H	Q3h
791	Q1h	CF3	H	Q3i
792	Q1h	CF3	H	Q3j
793	Q1h	CF3	H	Q3k
794	Q1h	CF3	H	Q3l
795	Q1h	CF3	H	Q3m
796	Q1h	CF3	H	Q3n
797	Q1h	CF3	H	Q3o
798	Q1h	CF3	H	Q3p
799	Q1h	CF3	H	Q3q
800	Q1h	CF3	Me	Q3a
801	Q1h	CF3	Me	Q3b
802	Q1h	CF3	Me	Q3c
803	Q1h	CF3	Me	Q3d
804	Q1h	CF3	Me	Q3e
805	Q1h	CF3	Me	Q3f
806	Q1h	CF3	Me	Q3g
807	Q1h	CF3	Me	Q3h
808	Q1h	CF3	Me	Q3i
809	Q1h	CF3	Me	Q3j
810	Q1h	CF3	Me	Q3k
811	Q1h	CF3	Me	Q3l
812	Q1h	CF3	Me	Q3m
813	Q1h	CF3	Me	Q3n
814	Q1h	CF3	Me	Q3o
815	Q1h	CF3	Me	Q3p
816	Q1h	CF3	Me	Q3q
817	Q1i	H	H	Q3a
818	Q1i	H	H	Q3b
819	Q1i	H	H	Q3c
820	Q1i	H	H	Q3d
821	Q1i	H	H	Q3e
822	Q1i	H	H	Q3f
823	Q1i	H	H	Q3g
824	Q1i	H	H	Q3h
825	Q1i	H	H	Q3i

826	Q1i	H	H	Q3j
827	Q1i	H	H	Q3k
828	Q1i	H	H	Q3l
829	Q1i	H	H	Q3m
830	Q1i	H	H	Q3n
831	Q1i	H	H	Q3o
832	Q1i	H	H	Q3p
833	Q1i	H	H	Q3q
834	Q1i	H	Me	Q3a
835	Q1i	H	Me	Q3b
836	Q1i	H	Me	Q3c
837	Q1i	H	Me	Q3d
838	Q1i	H	Me	Q3e
839	Q1i	H	Me	Q3f
840	Q1i	H	Me	Q3g
841	Q1i	H	Me	Q3h
842	Q1i	H	Me	Q3i
843	Q1i	H	Me	Q3j
844	Q1i	H	Me	Q3k
845	Q1i	H	Me	Q3l
846	Q1i	H	Me	Q3m
847	Q1i	H	Me	Q3n
848	Q1i	H	Me	Q3o
849	Q1i	H	Me	Q3p
850	Q1i	H	Me	Q3q
851	Q1i	Me	H	Q3a
852	Q1i	Me	H	Q3b
853	Q1i	Me	H	Q3c
854	Q1i	Me	H	Q3d
855	Q1i	Me	H	Q3e
856	Q1i	Me	H	Q3f
857	Q1i	Me	H	Q3g
858	Q1i	Me	H	Q3h
859	Q1i	Me	H	Q3i
860	Q1i	Me	H	Q3j
861	Q1i	Me	H	Q3k

862	Q1i	Me	H	Q3l
863	Q1i	Me	H	Q3m
864	Q1i	Me	H	Q3n
865	Q1i	Me	H	Q3o
866	Q1i	Me	H	Q3p
867	Q1i	Me	H	Q3q
868	Q1i	Me	Me	Q3a
869	Q1i	Me	Me	Q3b
870	Q1i	Me	Me	Q3c
871	Q1i	Me	Me	Q3d
872	Q1i	Me	Me	Q3e
873	Q1i	Me	Me	Q3f
874	Q1i	Me	Me	Q3g
875	Q1i	Me	Me	Q3h
876	Q1i	Me	Me	Q3i
877	Q1i	Me	Me	Q3j
878	Q1i	Me	Me	Q3k
879	Q1i	Me	Me	Q3l
880	Q1i	Me	Me	Q3m
881	Q1i	Me	Me	Q3n
882	Q1i	Me	Me	Q3o
883	Q1i	Me	Me	Q3p
884	Q1i	Me	Me	Q3q
885	Q1i	CF3	H	Q3a
886	Q1i	CF3	H	Q3b
887	Q1i	CF3	H	Q3c
888	Q1i	CF3	H	Q3d
889	Q1i	CF3	H	Q3e
890	Q1i	CF3	H	Q3f
891	Q1i	CF3	H	Q3g
892	Q1i	CF3	H	Q3h
893	Q1i	CF3	H	Q3i
894	Q1i	CF3	H	Q3j
895	Q1i	CF3	H	Q3k
896	Q1i	CF3	H	Q3l
897	Q1i	CF3	H	Q3m

898	Q1i	CF3	H	Q3n
899	Q1i	CF3	H	Q3o
900	Q1i	CF3	H	Q3p
901	Q1i	CF3	H	Q3q
902	Q1i	CF3	Me	Q3a
903	Q1i	CF3	Me	Q3b
904	Q1i	CF3	Me	Q3c
905	Q1i	CF3	Me	Q3d
906	Q1i	CF3	Me	Q3e
907	Q1i	CF3	Me	Q3f
908	Q1i	CF3	Me	Q3g
909	Q1i	CF3	Me	Q3h
910	Q1i	CF3	Me	Q3i
911	Q1i	CF3	Me	Q3j
912	Q1i	CF3	Me	Q3k
913	Q1i	CF3	Me	Q3l
914	Q1i	CF3	Me	Q3m
915	Q1i	CF3	Me	Q3n
916	Q1i	CF3	Me	Q3o
917	Q1i	CF3	Me	Q3p
918	Q1i	CF3	Me	Q3q
919	Q1j	H	H	Q3a
920	Q1j	H	H	Q3b
921	Q1j	H	H	Q3c
922	Q1j	H	H	Q3d
923	Q1j	H	H	Q3e
924	Q1j	H	H	Q3f
925	Q1j	H	H	Q3g
926	Q1j	H	H	Q3h
927	Q1j	H	H	Q3i
928	Q1j	H	H	Q3j
929	Q1j	H	H	Q3k
930	Q1j	H	H	Q3l
931	Q1j	H	H	Q3m
932	Q1j	H	H	Q3n
933	Q1j	H	H	Q3o

934	Q1j	H	H	Q3p
935	Q1j	H	H	Q3q
936	Q1j	H	Me	Q3a
937	Q1j	H	Me	Q3b
938	Q1j	H	Me	Q3c
939	Q1j	H	Me	Q3d
940	Q1j	H	Me	Q3e
941	Q1j	H	Me	Q3f
942	Q1j	H	Me	Q3g
943	Q1j	H	Me	Q3h
944	Q1j	H	Me	Q3i
945	Q1j	H	Me	Q3j
946	Q1j	H	Me	Q3k
947	Q1j	H	Me	Q3l
948	Q1j	H	Me	Q3m
949	Q1j	H	Me	Q3n
950	Q1j	H	Me	Q3o
951	Q1j	H	Me	Q3p
952	Q1j	H	Me	Q3q
953	Q1j	Me	H	Q3a
954	Q1j	Me	H	Q3b
955	Q1j	Me	H	Q3c
956	Q1j	Me	H	Q3d
957	Q1j	Me	H	Q3e
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959	Q1j	Me	H	Q3g
960	Q1j	Me	H	Q3h
961	Q1j	Me	H	Q3i
962	Q1j	Me	H	Q3j
963	Q1j	Me	H	Q3k
964	Q1j	Me	H	Q3l
965	Q1j	Me	H	Q3m
966	Q1j	Me	H	Q3n
967	Q1j	Me	H	Q3o
968	Q1j	Me	H	Q3p
969	Q1j	Me	H	Q3q

970	Q1j	Me	Me	Q3a
971	Q1j	Me	Me	Q3b
972	Q1j	Me	Me	Q3c
973	Q1j	Me	Me	Q3d
974	Q1j	Me	Me	Q3e
975	Q1j	Me	Me	Q3f
976	Q1j	Me	Me	Q3g
977	Q1j	Me	Me	Q3h
978	Q1j	Me	Me	Q3i
979	Q1j	Me	Me	Q3j
980	Q1j	Me	Me	Q3k
981	Q1j	Me	Me	Q3l
982	Q1j	Me	Me	Q3m
983	Q1j	Me	Me	Q3n
984	Q1j	Me	Me	Q3o
985	Q1j	Me	Me	Q3p
986	Q1j	Me	Me	Q3q
987	Q1j	CF3	H	Q3a
988	Q1j	CF3	H	Q3b
989	Q1j	CF3	H	Q3c
990	Q1j	CF3	H	Q3d
991	Q1j	CF3	H	Q3e
992	Q1j	CF3	H	Q3f
993	Q1j	CF3	H	Q3g
994	Q1j	CF3	H	Q3h
995	Q1j	CF3	H	Q3i
996	Q1j	CF3	H	Q3j
997	Q1j	CF3	H	Q3k
998	Q1j	CF3	H	Q3l
999	Q1j	CF3	H	Q3m
1000	Q1j	CF3	H	Q3n
1001	Q1j	CF3	H	Q3o
1002	Q1j	CF3	H	Q3p
1003	Q1j	CF3	H	Q3q
1004	Q1j	CF3	Me	Q3a
1005	Q1j	CF3	Me	Q3b

1006	Q1j	CF3	Me	Q3c
1007	Q1j	CF3	Me	Q3d
1008	Q1j	CF3	Me	Q3e
1009	Q1j	CF3	Me	Q3f
1010	Q1j	CF3	Me	Q3g
1011	Q1j	CF3	Me	Q3h
1012	Q1j	CF3	Me	Q3i
1013	Q1j	CF3	Me	Q3j
1014	Q1j	CF3	Me	Q3k
1015	Q1j	CF3	Me	Q3l
1016	Q1j	CF3	Me	Q3m
1017	Q1j	CF3	Me	Q3n
1018	Q1j	CF3	Me	Q3o
1019	Q1j	CF3	Me	Q3p
1020	Q1j	CF3	Me	Q3q
1021	Q1k	H	H	Q3a
1022	Q1k	H	H	Q3b
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1024	Q1k	H	H	Q3d
1025	Q1k	H	H	Q3e
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1027	Q1k	H	H	Q3g
1028	Q1k	H	H	Q3h
1029	Q1k	H	H	Q3i
1030	Q1k	H	H	Q3j
1031	Q1k	H	H	Q3k
1032	Q1k	H	H	Q3l
1033	Q1k	H	H	Q3m
1034	Q1k	H	H	Q3n
1035	Q1k	H	H	Q3o
1036	Q1k	H	H	Q3p
1037	Q1k	H	H	Q3q
1038	Q1k	H	Me	Q3a
1039	Q1k	H	Me	Q3b
1040	Q1k	H	Me	Q3c
1041	Q1k	H	Me	Q3d

1042	Q1k	H	Me	Q3e
1043	Q1k	H	Me	Q3f
1044	Q1k	H	Me	Q3g
1045	Q1k	H	Me	Q3h
1046	Q1k	H	Me	Q3i
1047	Q1k	H	Me	Q3j
1048	Q1k	H	Me	Q3k
1049	Q1k			

1065	Q1k	Me	H	Q3k
1066	Q1k	Me	H	Q3l
1067	Q1k	Me	H	Q3m
1068	Q1k	Me	H	Q3n
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1070	Q1k	Me	H	Q3p
1071	Q1k	Me	H	Q3q
1072	Q1k	Me	Me	Q3a
1073	Q1k	Me	Me	Q3b
1074	Q1k	Me	Me	Q3c
1075	Q1k	Me	Me	Q3d
1076	Q1k	Me	Me	Q3e
1077	Q1k	Me	Me	Q3f
1078	Q1k	Me	Me	Q3g
1079	Q1k	Me	Me	Q3h
1080	Q1k	Me	Me	Q3i
1081	Q1k	Me	Me	Q3j
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1083	Q1k	Me	Me	Q3l
1084	Q1k	Me	Me	Q3m
1085	Q1k	Me	Me	Q3n
1086	Q1k	Me	Me	Q3o
1087	Q1k	Me	Me	Q3p
1088	Q1k	Me	Me	Q3q
1089	Q1k	CF3	H	Q3a
1090	Q1k	CF3	H	Q3b
1091	Q1k	CF3	H	Q3c
1092	Q1k	CF3	H	Q3d
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1095	Q1k	CF3	H	Q3g
1096	Q1k	CF3	H	Q3h
1097	Q1k	CF3	H	Q3i
1098	Q1k	CF3	H	Q3j
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1100	Q1k	CF3	H	Q3l

1101	Q1k	CF3	H	Q3m
1102	Q1k	CF3	H	Q3n
1103	Q1k	CF3	H	Q3o
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1105	Q1k	CF3	H	Q3q
1106	Q1k	CF3	Me	Q3a
1107	Q1k	CF3	Me	Q3b
1108	Q1k	CF3	Me	Q3c
1109	Q1k	CF3	Me	Q3d
1110	Q1k	CF3	Me	Q3e
1111	Q1k	CF3	Me	Q3f
1112	Q1k	CF3	Me	Q3g
1113	Q1k	CF3	Me	Q3h
1114	Q1k	CF3	Me	Q3i
1115	Q1k	CF3	Me	Q3j
1116	Q1k	CF3	Me	Q3k
1117	Q1k	CF3	Me	Q3l
1118	Q1k	CF3	Me	Q3m
1119	Q1k	CF3	Me	Q3n
1120	Q1k	CF3	Me	Q3o
1121	Q1k	CF3	Me	Q3p
1122	Q1k	CF3	Me	Q3q
1123	Q1l	H	H	Q3a
1124	Q1l	H	H	Q3b
1125	Q1l	H	H	Q3c
1126	Q1l	H	H	Q3d
1127	Q1l	H	H	Q3e
1128	Q1l	H	H	Q3f
1129	Q1l	H	H	Q3g
1130	Q1l	H	H	Q3h
1131	Q1l	H	H	Q3i
1132	Q1l	H	H	Q3j
1133	Q1l	H	H	Q3k
1134	Q1l	H	H	Q3l
1135	Q1l	H	H	Q3m
1136	Q1l	H	H	Q3n

1137	Q11	H	H	Q3o
1138	Q11	H	H	Q3p
1139	Q11	H	H	Q3q
1140	Q11	H	Me	Q3a
1141	Q11	H	Me	Q3b
1142	Q11	H	Me	Q3c
1143	Q11	H	Me	Q3d
1144	Q11	H	Me	Q3e
1145	Q11	H	Me	Q3f
1146	Q11	H	Me	Q3g
1147	Q11	H	Me	Q3h
1148	Q11	H	Me	Q3i
1149	Q11	H	Me	Q3j
1150	Q11	H	Me	Q3k
1151	Q11	H	Me	Q3l
1152	Q11	H	Me	Q3m
1153	Q11	H	Me	Q3n
1154	Q11	H	Me	Q3o
1155	Q11	H	Me	Q3p
1156	Q11	H	Me	Q3q
1157	Q11	Me	H	Q3a
1158	Q11	Me	H	Q3b
1159	Q11	Me	H	Q3c
1160	Q11	Me	H	Q3d
1161	Q11	Me	H	Q3e
1162	Q11	Me	H	Q3f
1163	Q11	Me	H	Q3g
1164	Q11	Me	H	Q3h
1165	Q11	Me	H	Q3i
1166	Q11	Me	H	Q3j
1167	Q11	Me	H	Q3k
1168	Q11	Me	H	Q3l
1169	Q11	Me	H	Q3m
1170	Q11	Me	H	Q3n
1171	Q11	Me	H	Q3o
1172	Q11	Me	H	Q3p

1173	Q11	Me	H	Q3q
1174	Q11	Me	Me	Q3a
1175	Q11	Me	Me	Q3b
1176	Q11	Me	Me	Q3c
1177	Q11	Me	Me	Q3d
1178	Q11	Me	Me	Q3e
1179	Q11	Me	Me	Q3f
1180	Q11	Me	Me	Q3g
1181	Q11	Me	Me	Q3h
1182	Q11	Me	Me	Q3i
1183	Q11	Me	Me	Q3j
1184	Q11	Me	Me	Q3k
1185	Q11	Me	Me	Q3l
1186	Q11	Me	Me	Q3m
1187	Q11	Me	Me	Q3n
1188	Q11	Me	Me	Q3o
1189	Q11	Me	Me	Q3p
1190	Q11	Me	Me	Q3q
1191	Q11	CF3	H	Q3a
1192	Q11	CF3	H	Q3b
1193	Q11	CF3	H	Q3c
1194	Q11	CF3	H	Q3d
1195	Q11	CF3	H	Q3e
1196	Q11	CF3	H	Q3f
1197	Q11	CF3	H	Q3g
1198	Q11	CF3	H	Q3h
1199	Q11	CF3	H	Q3i
1200	Q11	CF3	H	Q3j
1201	Q11	CF3	H	Q3k
1202	Q11	CF3	H	Q3l
1203	Q11	CF3	H	Q3m
1204	Q11	CF3	H	Q3n
1205	Q11	CF3	H	Q3o
1206	Q11	CF3	H	Q3p
1207	Q11	CF3	H	Q3q
1208	Q11	CF3	Me	Q3a

1209	Q1l	CF3	Me	Q3b
1210	Q1l	CF3	Me	Q3c
1211	Q1l	CF3	Me	Q3d
1212	Q1l	CF3	Me	Q3e
1213	Q1l	CF3	Me	Q3f
1214	Q1l	CF3	Me	Q3g
1215	Q1l	CF3	Me	Q3h
1216	Q1l	CF3	Me	Q3i
1217	Q1l	CF3	Me	Q3j
1218	Q1l	CF3	Me	Q3k
1219	Q1l	CF3	Me	Q3l
1220	Q1l	CF3	Me	Q3m
1221	Q1l	CF3	Me	Q3n
1222	Q1l	CF3	Me	Q3o
1223	Q1l	CF3	Me	Q3p
1224	Q1l	CF3	Me	Q3q
1225	Q1m	H	H	Q3a
1226	Q1m	H	H	Q3b
1227	Q1m	H	H	Q3c
1228	Q1m	H	H	Q3d
1229	Q1m	H	H	Q3e
1230	Q1m	H	H	Q3f
1231	Q1m	H	H	Q3g
1232	Q1m	H	H	Q3h
1233	Q1m	H	H	Q3i
1234	Q1m	H	H	Q3j
1235	Q1m	H	H	Q3k
1236	Q1m	H	H	Q3l
1237	Q1m	H	H	Q3m
1238	Q1m	H	H	Q3n
1239	Q1m	H	H	Q3o
1240	Q1m	H	H	Q3p
1241	Q1m	H	H	Q3q
1242	Q1m	H	Me	Q3a
1243	Q1m	H	Me	Q3b
1244	Q1m	H	Me	Q3c

1245	Q1m	H	Me	Q3d
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1247	Q1m	H	Me	Q3f
1248	Q1m	H	Me	Q3g
1249	Q1m	H	Me	Q3h
1250	Q1m	H	Me	Q3i
1251	Q1m	H	Me	Q3j
1252	Q1m	H	Me	Q3k
1253	Q1m	H	Me	Q3l
1254	Q1m	H	Me	Q3m
1255	Q1m	H	Me	Q3n
1256	Q1m	H	Me	Q3o
1257	Q1m	H	Me	Q3p
1258	Q1m	H	Me	Q3q
1259	Q1m	Me	H	Q3a
1260	Q1m	Me	H	Q3b
1261	Q1m	Me	H	Q3c
1262	Q1m	Me	H	Q3d
1263	Q1m	Me	H	Q3e
1264	Q1m	Me	H	Q3f
1265	Q1m	Me	H	Q3g
1266	Q1m	Me	H	Q3h
1267	Q1m	Me	H	Q3i
1268	Q1m	Me	H	Q3j
1269	Q1m	Me	H	Q3k
1270	Q1m	Me	H	Q3l
1271	Q1m	Me	H	Q3m
1272	Q1m	Me	H	Q3n
1273	Q1m	Me	H	Q3o
1274	Q1m	Me	H	Q3p
1275	Q1m	Me	H	Q3q
1276	Q1m	Me	Me	Q3a
1277	Q1m	Me	Me	Q3b
1278	Q1m	Me	Me	Q3c
1279	Q1m	Me	Me	Q3d
1280	Q1m	Me	Me	Q3e

1281	Q1m	Me	Me	Q3f
1282	Q1m	Me	Me	Q3g
1283	Q1m	Me	Me	Q3h
1284	Q1m	Me	Me	Q3i
1285	Q1m	Me	Me	Q3j
1286	Q1m	Me	Me	Q3k
1287	Q1m	Me	Me	Q3l
1288	Q1m	Me	Me	Q3m
1289	Q1m	Me	Me	Q3n
1290	Q1m	Me	Me	Q3o
1291	Q1m	Me	Me	Q3p
1292	Q1m	Me	Me	Q3q
1293	Q1m	CF3	H	Q3a
1294	Q1m	CF3	H	Q3b
1295	Q1m	CF3	H	Q3c
1296	Q1m	CF3	H	Q3d
1297	Q1m	CF3	H	Q3e
1298	Q1m	CF3	H	Q3f
1299	Q1m	CF3	H	Q3g
1300	Q1m	CF3	H	Q3h
1301	Q1m	CF3	H	Q3i
1302	Q1m	CF3	H	Q3j
1303	Q1m	CF3	H	Q3k
1304	Q1m	CF3	H	Q3l
1305	Q1m	CF3	H	Q3m
1306	Q1m	CF3	H	Q3n
1307	Q1m	CF3	H	Q3o
1308	Q1m	CF3	H	Q3p
1309	Q1m	CF3	H	Q3q
1310	Q1m	CF3	Me	Q3a
1311	Q1m	CF3	Me	Q3b
1312	Q1m	CF3	Me	Q3c
1313	Q1m	CF3	Me	Q3d
1314	Q1m	CF3	Me	Q3e
1315	Q1m	CF3	Me	Q3f
1316	Q1m	CF3	Me	Q3g

1317	Q1m	CF3	Me	Q3h
1318	Q1m	CF3	Me	Q3i
1319	Q1m	CF3	Me	Q3j
1320	Q1m	CF3	Me	Q3k
1321	Q1m	CF3	Me	Q3l
1322	Q1m	CF3	Me	Q3m
1323	Q1m	CF3	Me	Q3n
1324	Q1m	CF3	Me	Q3o
1325	Q1m	CF3	Me	Q3p
1326	Q1m	CF3	Me	Q3q
1327	Q1n	H	H	Q3a
1328	Q1n	H	H	Q3b
1329	Q1n	H	H	Q3c
1330	Q1n	H	H	Q3d
1331	Q1n	H	H	Q3e
1332	Q1n	H	H	Q3f
1333	Q1n	H	H	Q3g
1334	Q1n	H	H	Q3h
1335	Q1n	H	H	Q3i
1336	Q1n	H	H	Q3j
1337	Q1n	H	H	Q3k
1338	Q1n	H	H	Q3l
1339	Q1n	H	H	Q3m
1340	Q1n	H	H	Q3n
1341	Q1n	H	H	Q3o
1342	Q1n	H	H	Q3p
1343	Q1n	H	H	Q3q
1344	Q1n	H	Me	Q3a
1345	Q1n	H	Me	Q3b
1346	Q1n	H	Me	Q3c
1347	Q1n	H	Me	Q3d
1348	Q1n	H	Me	Q3e
1349	Q1n	H	Me	Q3f
1350	Q1n	H	Me	Q3g
1351	Q1n	H	Me	Q3h
1352	Q1n	H	Me	Q3i

1353	Q1n	H	Me	Q3j
1354	Q1n	H	Me	Q3k
1355	Q1n	H	Me	Q3l
1356	Q1n	H	Me	Q3m
1357	Q1n	H	Me	Q3n
1358	Q1n	H	Me	Q3o
1359	Q1n	H	Me	Q3p
1360	Q1n	H	Me	Q3q
1361	Q1n	Me	H	Q3a
1362	Q1n	Me	H	Q3b
1363	Q1n	Me	H	Q3c
1364	Q1n	Me	H	Q3d
1365	Q1n	Me	H	Q3e
1366	Q1n	Me	H	Q3f
1367	Q1n	Me	H	Q3g
1368	Q1n	Me	H	Q3h
1369	Q1n	Me	H	Q3i
1370	Q1n	Me	H	Q3j
1371	Q1n	Me	H	Q3k
1372	Q1n	Me	H	Q3l
1373	Q1n	Me	H	Q3m
1374	Q1n	Me	H	Q3n
1375	Q1n	Me	H	Q3o
1376	Q1n	Me	H	Q3p
1377	Q1n	Me	H	Q3q
1378	Q1n	Me	Me	Q3a
1379	Q1n	Me	Me	Q3b
1380	Q1n	Me	Me	Q3c
1381	Q1n	Me	Me	Q3d
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1384	Q1n	Me	Me	Q3g
1385	Q1n	Me	Me	Q3h
1386	Q1n	Me	Me	Q3i
1387	Q1n	Me	Me	Q3j
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1389	Q1n	Me	Me	Q3l
1390	Q1n	Me	Me	Q3m
1391	Q1n	Me	Me	Q3n
1392	Q1n	Me	Me	Q3o
1393	Q1n	Me	Me	Q3p
1394	Q1n	Me	Me	Q3q
1395	Q1n	CF3	H	Q3a
1396	Q1n	CF3	H	Q3b
1397	Q1n	CF3	H	Q3c
1398	Q1n	CF3	H	Q3d
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1409	Q1n	CF3	H	Q3o
1410	Q1n	CF3	H	Q3p
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1422	Q1n	CF3	Me	Q3k
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1424	Q1n	CF3	Me	Q3m

1425	Q1n	CF3	Me	Q3n
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1427	Q1n	CF3	Me	Q3p
1428	Q1n	CF3	Me	Q3q
1429	Q1o	H	H	Q3a
1430	Q1o	H	H	Q3b
1431	Q1o	H	H	Q3c
1432	Q1o	H	H	Q3d
1433	Q1o	H	H	Q3e
1434	Q1o	H	H	Q3f
1435	Q1o	H	H	Q3g
1436	Q1o	H	H	Q3h
1437	Q1o	H	H	Q3i
1438	Q1o	H	H	Q3j
1439	Q1o	H	H	Q3k
1440	Q1o	H	H	Q3l
1441	Q1o	H	H	Q3m
1442	Q1o	H	H	Q3n
1443	Q1o	H	H	Q3o
1444	Q1o	H	H	Q3p
1445	Q1o	H	H	Q3q
1446	Q1o	H	Me	Q3a
1447	Q1o	H	Me	Q3b
1448	Q1o	H	Me	Q3c
1449	Q1o	H	Me	Q3d
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1452	Q1o	H	Me	Q3g
1453	Q1o	H	Me	Q3h
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1457	Q1o	H	Me	Q3l
1458	Q1o	H	Me	Q3m
1459	Q1o	H	Me	Q3n
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1461	Q1o	H	Me	Q3p
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1463	Q1o	Me	H	Q3a
1464	Q1o	Me	H	Q3b
1465	Q1o	Me	H	Q3c
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1470	Q1o	Me	H	Q3h
1471	Q1o	Me	H	Q3i
1472	Q1o	Me	H	Q3j
1473	Q1o	Me	H	Q3k
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1481	Q1o	Me	Me	Q3b
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1486	Q1o	Me	Me	Q3g
1487	Q1o	Me	Me	Q3h
1488	Q1o	Me	Me	Q3i
1489	Q1o	Me	Me	Q3j
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1491	Q1o	Me	Me	Q3l
1492	Q1o	Me	Me	Q3m
1493	Q1o	Me	Me	Q3n
1494	Q1o	Me	Me	Q3o
1495	Q1o	Me	Me	Q3p
1496	Q1o	Me	Me	Q3q

1497	Q1o	CF3	H	Q3a
1498	Q1o	CF3	H	Q3b
1499	Q1o	CF3	H	Q3c
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1501	Q1o	CF3	H	Q3e
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1503	Q1o	CF3	H	Q3g
1504	Q1o	CF3	H	Q3h
1505	Q1o	CF3	H	Q3i
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1527	Q1o	CF3	Me	Q3n
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1531	Q1p	H	H	Q3a
1532	Q1p	H	H	Q3b

1533	Q1p	H	H	Q3c
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1535	Q1p	H	H	Q3e
1536	Q1p	H	H	Q3f
1537	Q1p	H	H	Q3g
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1549	Q1p	H	Me	Q3b
1550	Q1p	H	Me	Q3c
1551	Q1p	H	Me	Q3d
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1554	Q1p	H	Me	Q3g
1555	Q1p	H	Me	Q3h
1556	Q1p	H	Me	Q3i
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1559	Q1p	H	Me	Q3l
1560	Q1p	H	Me	Q3m
1561	Q1p	H	Me	Q3n
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1564	Q1p	H	Me	Q3q
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1566	Q1p	Me	H	Q3b
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1581	Q1p	Me	H	Q3q
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1583	Q1p	Me	Me	Q3b
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1596	Q1p	Me	Me	Q3o
1597	Q1p	Me	Me	Q3p
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1599	Q1p	CF3	H	Q3a
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1601	Q1p	CF3	H	Q3c
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1614	Q1p	CF3	H	Q3p
1615	Q1p	CF3	H	Q3q
1616	Q1p	CF3	Me	Q3a
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1623	Q1p	CF3	Me	Q3h
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1625	Q1p	CF3	Me	Q3j
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1627	Q1p	CF3	Me	Q3l
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1630	Q1p	CF3	Me	Q3o
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1635	Q1q	H	H	Q3c
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1639	Q1q	H	H	Q3g
1640	Q1q	H	H	Q3h

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1649	Q1q	H	H	Q3q
1650	Q1q	H	Me	Q3a
1651	Q1q	H	Me	Q3b
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1653	Q1q	H	Me	Q3d
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1663	Q1q	H	Me	Q3n
1664	Q1q	H	Me	Q3o
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1674	Q1q	Me	H	Q3h
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1732	Q1q	CF3	Me	Q3o
1733	Q1q	CF3	Me	Q3p
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1736	Q1r	H	H	Q3b
1737	Q1r	H	H	Q3c
1738	Q1r	H	H	Q3d
1739	Q1r	H	H	Q3e
1740	Q1r	H	H	Q3f
1741	Q1r	H	H	Q3g
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1746	Q1r	H	H	Q3l
1747	Q1r	H	H	Q3m
1748	Q1r	H	H	Q3n

1749	Q1r	H	H	Q3o
1750	Q1r	H	H	Q3p
1751	Q1r	H	H	Q3q
1752	Q1r	H	Me	Q3a
1753	Q1r	H	Me	Q3b
1754	Q1r	H	Me	Q3c
1755	Q1r	H	Me	Q3d
1756	Q1r	H	Me	Q3e
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1761	Q1r	H	Me	Q3j
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1764	Q1r	H	Me	Q3m
1765	Q1r	H	Me	Q3n
1766	Q1r	H	Me	Q3o
1767	Q1r	H	Me	Q3p
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1770	Q1r	Me	H	Q3b
1771	Q1r	Me	H	Q3c
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1776	Q1r	Me	H	Q3h
1777	Q1r	Me	H	Q3i
1778	Q1r	Me	H	Q3j
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1780	Q1r	Me	H	Q3l
1781	Q1r	Me	H	Q3m
1782	Q1r	Me	H	Q3n
1783	Q1r	Me	H	Q3o
1784	Q1r	Me	H	Q3p

1785	Q1r	Me	H	Q3q
1786	Q1r	Me	Me	Q3a
1787	Q1r	Me	Me	Q3b
1788	Q1r	Me	Me	Q3c
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1792	Q1r	Me	Me	Q3g
1793	Q1r	Me	Me	Q3h
1794	Q1r	Me	Me	Q3i
1795	Q1r	Me	Me	Q3j
1796	Q1r	Me	Me	Q3k
1797	Q1r	Me	Me	Q3l
1798	Q1r	Me	Me	Q3m
1799	Q1r	Me	Me	Q3n
1800	Q1r	Me	Me	Q3o
1801	Q1r	Me	Me	Q3p
1802	Q1r	Me	Me	Q3q
1803	Q1r	CF3	H	Q3a
1804	Q1r	CF3	H	Q3b
1805	Q1r	CF3	H	Q3c
1806	Q1r	CF3	H	Q3d
1807	Q1r	CF3	H	Q3e
1808	Q1r	CF3	H	Q3f
1809	Q1r	CF3	H	Q3g
1810	Q1r	CF3	H	Q3h
1811	Q1r	CF3	H	Q3i
1812	Q1r	CF3	H	Q3j
1813	Q1r	CF3	H	Q3k
1814	Q1r	CF3	H	Q3l
1815	Q1r	CF3	H	Q3m
1816	Q1r	CF3	H	Q3n
1817	Q1r	CF3	H	Q3o
1818	Q1r	CF3	H	Q3p
1819	Q1r	CF3	H	Q3q
1820	Q1r	CF3	Me	Q3a

1821	Q1r	CF3	Me	Q3b
1822	Q1r	CF3	Me	Q3c
1823	Q1r	CF3	Me	Q3d
1824	Q1r	CF3	Me	Q3e
1825	Q1r	CF3	Me	Q3f
1826	Q1r	CF3	Me	Q3g
1827	Q1r	CF3	Me	Q3h
1828	Q1r	CF3	Me	Q3i
1829	Q1r	CF3	Me	Q3j
1830	Q1r	CF3	Me	Q3k
1831	Q1r	CF3	Me	Q3l
1832	Q1r	CF3	Me	Q3m
1833	Q1r	CF3	Me	Q3n
1834	Q1r	CF3	Me	Q3o
1835	Q1r	CF3	Me	Q3p
1836	Q1r	CF3	Me	Q3q
1837	Q1s	H	H	Q3a
1838	Q1s	H	H	Q3b
1839	Q1s	H	H	Q3c
1840	Q1s	H	H	Q3d
1841	Q1s	H	H	Q3e
1842	Q1s	H	H	Q3f
1843	Q1s	H	H	Q3g
1844	Q1s	H	H	Q3h
1845	Q1s	H	H	Q3i
1846	Q1s	H	H	Q3j
1847	Q1s	H	H	Q3k
1848	Q1s	H	H	Q3l
1849	Q1s	H	H	Q3m
1850	Q1s	H	H	Q3n
1851	Q1s	H	H	Q3o
1852	Q1s	H	H	Q3p
1853	Q1s	H	H	Q3q
1854	Q1s	H	Me	Q3a
1855	Q1s	H	Me	Q3b
1856	Q1s	H	Me	Q3c

1857	Q1s	H	Me	Q3d
1858	Q1s	H	Me	Q3e
1859	Q1s	H	Me	Q3f
1860	Q1s	H	Me	Q3g
1861	Q1s	H	Me	Q3h
1862	Q1s	H	Me	Q3i
1863	Q1s	H	Me	Q3j
1864	Q1s	H	Me	Q3k
1865	Q1s	H	Me	Q3l
1866	Q1s	H	Me	Q3m
1867	Q1s	H	Me	Q3n
1868	Q1s	H	Me	Q3o
1869	Q1s	H	Me	Q3p
1870	Q1s	H	Me	Q3q
1871	Q1s	Me	H	Q3a
1872	Q1s	Me	H	Q3b
1873	Q1s	Me	H	Q3c
1874	Q1s	Me	H	Q3d
1875	Q1s	Me	H	Q3e
1876	Q1s	Me	H	Q3f
1877	Q1s	Me	H	Q3g
1878	Q1s	Me	H	Q3h
1879	Q1s	Me	H	Q3i
1880	Q1s	Me	H	Q3j
1881	Q1s	Me	H	Q3k
1882	Q1s	Me	H	Q3l
1883	Q1s	Me	H	Q3m
1884	Q1s	Me	H	Q3n
1885	Q1s	Me	H	Q3o
1886	Q1s	Me	H	Q3p
1887	Q1s	Me	H	Q3q
1888	Q1s	Me	Me	Q3a
1889	Q1s	Me	Me	Q3b
1890	Q1s	Me	Me	Q3c
1891	Q1s	Me	Me	Q3d
1892	Q1s	Me	Me	Q3e

1893	Q1s	Me	Me	Q3f
1894	Q1s	Me	Me	Q3g
1895	Q1s	Me	Me	Q3h
1896	Q1s	Me	Me	Q3i
1897	Q1s	Me	Me	Q3j
1898	Q1s	Me	Me	Q3k
1899	Q1s	Me	Me	Q3l
1900	Q1s	Me	Me	Q3m
1901	Q1s	Me	Me	Q3n
1902	Q1s	Me	Me	Q3o
1903	Q1s	Me	Me	Q3p
1904	Q1s	Me	Me	Q3q
1905	Q1s	CF3	H	Q3a
1906	Q1s	CF3	H	Q3b
1907	Q1s	CF3	H	Q3c
1908	Q1s	CF3	H	Q3d
1909	Q1s	CF3	H	Q3e
1910	Q1s	CF3	H	Q3f
1911	Q1s	CF3	H	Q3g
1912	Q1s	CF3	H	Q3h
1913	Q1s	CF3	H	Q3i
1914	Q1s	CF3	H	Q3j
1915	Q1s	CF3	H	Q3k
1916	Q1s	CF3	H	Q3l
1917	Q1s	CF3	H	Q3m
1918	Q1s	CF3	H	Q3n
1919	Q1s	CF3	H	Q3o
1920	Q1s	CF3	H	Q3p
1921	Q1s	CF3	H	Q3q
1922	Q1s	CF3	Me	Q3a
1923	Q1s	CF3	Me	Q3b
1924	Q1s	CF3	Me	Q3c
1925	Q1s	CF3	Me	Q3d
1926	Q1s	CF3	Me	Q3e
1927	Q1s	CF3	Me	Q3f
1928	Q1s	CF3	Me	Q3g

1929	Q1s	CF3	Me	Q3h
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1931	Q1s	CF3	Me	Q3j
1932	Q1s	CF3	Me	Q3k
1933	Q1s	CF3	Me	Q3l
1934	Q1s	CF3	Me	Q3m
1935	Q1s	CF3	Me	Q3n
1936	Q1s	CF3	Me	Q3o
1937	Q1s	CF3	Me	Q3p
1938	Q1s	CF3	Me	Q3q
1939	Q1t	H	H	Q3a
1940	Q1t	H	H	Q3b
1941	Q1t	H	H	Q3c
1942	Q1t	H	H	Q3d
1943	Q1t	H	H	Q3e
1944	Q1t	H	H	Q3f
1945	Q1t	H	H	Q3g
1946	Q1t	H	H	Q3h
1947	Q1t	H	H	Q3i
1948	Q1t	H	H	Q3j
1949	Q1t	H	H	Q3k
1950	Q1t	H	H	Q3l
1951	Q1t	H	H	Q3m
1952	Q1t	H	H	Q3n
1953	Q1t	H	H	Q3o
1954	Q1t	H	H	Q3p
1955	Q1t	H	H	Q3q
1956	Q1t	H	Me	Q3a
1957	Q1t	H	Me	Q3b
1958	Q1t	H	Me	Q3c
1959	Q1t	H	Me	Q3d
1960	Q1t	H	Me	Q3e
1961	Q1t	H	Me	Q3f
1962	Q1t	H	Me	Q3g
1963	Q1t	H	Me	Q3h
1964	Q1t	H	Me	Q3i

1965	Q1t	H	Me	Q3j
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1967	Q1t	H	Me	Q3l
1968	Q1t	H	Me	Q3m
1969	Q1t	H	Me	Q3n
1970	Q1t	H	Me	Q3o
1971	Q1t	H	Me	Q3p
1972	Q1t	H	Me	Q3q
1973	Q1t	Me	H	Q3a
1974	Q1t	Me	H	Q3b
1975	Q1t	Me	H	Q3c
1976	Q1t	Me	H	Q3d
1977	Q1t	Me	H	Q3e
1978	Q1t	Me	H	Q3f
1979	Q1t	Me	H	Q3g
1980	Q1t	Me	H	Q3h
1981	Q1t	Me	H	Q3i
1982	Q1t	Me	H	Q3j
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1985	Q1t	Me	H	Q3m
1986	Q1t	Me	H	Q3n
1987	Q1t	Me	H	Q3o
1988	Q1t	Me	H	Q3p
1989	Q1t	Me	H	Q3q
1990	Q1t	Me	Me	Q3a
1991	Q1t	Me	Me	Q3b
1992	Q1t	Me	Me	Q3c
1993	Q1t	Me	Me	Q3d
1994	Q1t	Me	Me	Q3e
1995	Q1t	Me	Me	Q3f
1996	Q1t	Me	Me	Q3g
1997	Q1t	Me	Me	Q3h
1998	Q1t	Me	Me	Q3i
1999	Q1t	Me	Me	Q3j
2000	Q1t	Me	Me	Q3k

2001	Q1t	Me	Me	Q3l
2002	Q1t	Me	Me	Q3m
2003	Q1t	Me	Me	Q3n
2004	Q1t	Me	Me	Q3o
2005	Q1t	Me	Me	Q3p
2006	Q1t	Me	Me	Q3q
2007	Q1t	CF3	H	Q3a
2008	Q1t	CF3	H	Q3b
2009	Q1t	CF3	H	Q3c
2010	Q1t	CF3	H	Q3d
2011	Q1t	CF3	H	Q3e
2012	Q1t	CF3	H	Q3f
2013	Q1t	CF3	H	Q3g
2014	Q1t	CF3	H	Q3h
2015	Q1t	CF3	H	Q3i
2016	Q1t	CF3	H	Q3j
2017	Q1t	CF3	H	Q3k
2018	Q1t	CF3	H	Q3l
2019	Q1t	CF3	H	Q3m
2020	Q1t	CF3	H	Q3n
2021	Q1t	CF3	H	Q3o
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2023	Q1t	CF3	H	Q3q
2024	Q1t	CF3	Me	Q3a
2025	Q1t	CF3	Me	Q3b
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2039	Q1t	CF3	Me	Q3p
2040	Q1t	CF3	Me	Q3q
2041	Q1u	H	H	Q3a
2042	Q1u	H	H	Q3b
2043	Q1u	H	H	Q3c
2044	Q1u	H	H	Q3d
2045	Q1u	H	H	Q3e
2046	Q1u	H	H	Q3f
2047	Q1u	H	H	Q3g
2048	Q1u	H	H	Q3h
2049	Q1u	H	H	Q3i
2050	Q1u	H	H	Q3j
2051	Q1u	H	H	Q3k
2052	Q1u	H	H	Q3l
2053	Q1u	H	H	Q3m
2054	Q1u	H	H	Q3n
2055	Q1u	H	H	Q3o
2056	Q1u	H	H	Q3p
2057	Q1u	H	H	Q3q
2058	Q1u	H	Me	Q3a
2059	Q1u	H	Me	Q3b
2060	Q1u	H	Me	Q3c
2061	Q1u	H	Me	Q3d
2062	Q1u	H	Me	Q3e
2063	Q1u	H	Me	Q3f
2064	Q1u	H	Me	Q3g
2065	Q1u	H	Me	Q3h
2066	Q1u	H	Me	Q3i
2067	Q1u	H	Me	Q3j
2068	Q1u	H	Me	Q3k
2069	Q1u	H	Me	Q3l
2070	Q1u	H	Me	Q3m
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2073	Q1u	H	Me	Q3p
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2075	Q1u	Me	H	Q3a
2076	Q1u	Me	H	Q3b
2077	Q1u	Me	H	Q3c
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2080	Q1u	Me	H	Q3f
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2082	Q1u	Me	H	Q3h
2083	Q1u	Me	H	Q3i
2084	Q1u	Me	H	Q3j
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2087	Q1u	Me	H	Q3m
2088	Q1u	Me	H	Q3n
2089	Q1u	Me	H	Q3o
2090	Q1u	Me	H	Q3p
2091	Q1u	Me	H	Q3q
2092	Q1u	Me	Me	Q3a
2093	Q1u	Me	Me	Q3b
2094	Q1u	Me	Me	Q3c
2095	Q1u	Me	Me	Q3d
2096	Q1u	Me	Me	Q3e
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2098	Q1u	Me	Me	Q3g
2099	Q1u	Me	Me	Q3h
2100	Q1u	Me	Me	Q3i
2101	Q1u	Me	Me	Q3j
2102	Q1u	Me	Me	Q3k
2103	Q1u	Me	Me	Q3l
2104	Q1u	Me	Me	Q3m
2105	Q1u	Me	Me	Q3n
2106	Q1u	Me	Me	Q3o
2107	Q1u	Me	Me	Q3p
2108	Q1u	Me	Me	Q3q

2109	Q1u	CF3	H	Q3a
2110	Q1u	CF3	H	Q3b
2111	Q1u	CF3	H	Q3c
2112	Q1u	CF3	H	Q3d
2113	Q1u	CF3	H	Q3e
2114	Q1u	CF3	H	Q3f
2115	Q1u	CF3	H	Q3g
2116	Q1u	CF3	H	Q3h
2117	Q1u	CF3	H	Q3i
2118	Q1u	CF3	H	Q3j
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2122	Q1u	CF3	H	Q3n
2123	Q1u	CF3	H	Q3o
2124	Q1u	CF3	H	Q3p
2125	Q1u	CF3	H	Q3q
2126	Q1u	CF3	Me	Q3a
2127	Q1u	CF3	Me	Q3b
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2133	Q1u	CF3	Me	Q3h
2134	Q1u	CF3	Me	Q3i
2135	Q1u	CF3	Me	Q3j
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2137	Q1u	CF3	Me	Q3l
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2139	Q1u	CF3	Me	Q3n
2140	Q1u	CF3	Me	Q3o
2141	Q1u	CF3	Me	Q3p
2142	Q1u	CF3	Me	Q3q
2143	Q1v	H	H	Q3a
2144	Q1v	H	H	Q3b

2145	Q1v	H	H	Q3c
2146	Q1v	H	H	Q3d
2147	Q1v	H	H	Q3e
2148	Q1v	H	H	Q3f
2149	Q1v	H	H	Q3g
2150	Q1v	H	H	Q3h
2151	Q1v	H	H	Q3i
2152	Q1v	H	H	Q3j
2153	Q1v	H	H	Q3k
2154	Q1v	H	H	Q3l
2155	Q1v	H	H	Q3m
2156	Q1v	H	H	Q3n
2157	Q1v	H	H	Q3o
2158	Q1v	H	H	Q3p
2159	Q1v	H	H	Q3q
2160	Q1v	H	Me	Q3a
2161	Q1v	H	Me	Q3b
2162	Q1v	H	Me	Q3c
2163	Q1v	H	Me	Q3d
2164	Q1v	H	Me	Q3e
2165	Q1v	H	Me	Q3f
2166	Q1v	H	Me	Q3g
2167	Q1v	H	Me	Q3h
2168	Q1v	H	Me	Q3i
2169	Q1v	H	Me	Q3j
2170	Q1v	H	Me	Q3k
2171	Q1v	H	Me	Q3l
2172	Q1v	H	Me	Q3m
2173	Q1v	H	Me	Q3n
2174	Q1v	H	Me	Q3o
2175	Q1v	H	Me	Q3p
2176	Q1v	H	Me	Q3q
2177	Q1v	Me	H	Q3a
2178	Q1v	Me	H	Q3b
2179	Q1v	Me	H	Q3c
2180	Q1v	Me	H	Q3d

2181	Q1v	Me	H	Q3e
2182	Q1v	Me	H	Q3f
2183	Q1v	Me	H	Q3g
2184	Q1v	Me	H	Q3h
2185	Q1v	Me	H	Q3i
2186	Q1v	Me	H	Q3j
2187	Q1v	Me	H	Q3k
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2189	Q1v	Me	H	Q3m
2190	Q1v	Me	H	Q3n
2191	Q1v	Me	H	Q3o
2192	Q1v	Me	H	Q3p
2193	Q1v	Me	H	Q3q
2194	Q1v	Me	Me	Q3a
2195	Q1v	Me	Me	Q3b
2196	Q1v	Me	Me	Q3c
2197	Q1v	Me	Me	Q3d
2198	Q1v	Me	Me	Q3e
2199	Q1v	Me	Me	Q3f
2200	Q1v	Me	Me	Q3g
2201	Q1v	Me	Me	Q3h
2202	Q1v	Me	Me	Q3i
2203	Q1v	Me	Me	Q3j
2204	Q1v	Me	Me	Q3k
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2206	Q1v	Me	Me	Q3m
2207	Q1v	Me	Me	Q3n
2208	Q1v	Me	Me	Q3o
2209	Q1v	Me	Me	Q3p
2210	Q1v	Me	Me	Q3q
2211	Q1v	CF3	H	Q3a
2212	Q1v	CF3	H	Q3b
2213	Q1v	CF3	H	Q3c
2214	Q1v	CF3	H	Q3d
2215	Q1v	CF3	H	Q3e
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2219	Q1v	CF3	H	Q3i
2220	Q1v	CF3	H	Q3j
2221	Q1v	CF3	H	Q3k
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2224	Q1v	CF3	H	Q3n
2225	Q1v	CF3	H	Q3o
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2229	Q1v	CF3	Me	Q3b
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2233	Q1v	CF3	Me	Q3f
2234	Q1v	CF3	Me	Q3g
2235	Q1v	CF3	Me	Q3h
2236	Q1v	CF3	Me	Q3i
2237	Q1v	CF3	Me	Q3j
2238	Q1v	CF3	Me	Q3k
2239	Q1v	CF3	Me	Q3l
2240	Q1v	CF3	Me	Q3m
2241	Q1v	CF3	Me	Q3n
2242	Q1v	CF3	Me	Q3o
2243	Q1v	CF3	Me	Q3p
2244	Q1v	CF3	Me	Q3q
2245	Q1w	H	H	Q3a
2246	Q1w	H	H	Q3b
2247	Q1w	H	H	Q3c
2248	Q1w	H	H	Q3d
2249	Q1w	H	H	Q3e
2250	Q1w	H	H	Q3f
2251	Q1w	H	H	Q3g
2252	Q1w	H	H	Q3h

2253	Q1w	H	H	Q3i
2254	Q1w	H	H	Q3j
2255	Q1w	H	H	Q3k
2256	Q1w	H	H	Q3l
2257	Q1w	H	H	Q3m
2258	Q1w	H	H	Q3n
2259	Q1w	H	H	Q3o
2260	Q1w	H	H	Q3p
2261	Q1w	H	H	Q3q
2262	Q1w	H	Me	Q3a
2263	Q1w	H	Me	Q3b
2264	Q1w	H	Me	Q3c
2265	Q1w	H	Me	Q3d
2266	Q1w	H	Me	Q3e
2267	Q1w	H	Me	Q3f
2268	Q1w	H	Me	Q3g
2269	Q1w	H	Me	Q3h
2270	Q1w	H	Me	Q3i
2271	Q1w	H	Me	Q3j
2272	Q1w	H	Me	Q3k
2273	Q1w	H	Me	Q3l
2274	Q1w	H	Me	Q3m
2275	Q1w	H	Me	Q3n
2276	Q1w	H	Me	Q3o
2277	Q1w	H	Me	Q3p
2278	Q1w	H	Me	Q3q
2279	Q1w	Me	H	Q3a
2280	Q1w	Me	H	Q3b
2281	Q1w	Me	H	Q3c
2282	Q1w	Me	H	Q3d
2283	Q1w	Me	H	Q3e
2284	Q1w	Me	H	Q3f
2285	Q1w	Me	H	Q3g
2286	Q1w	Me	H	Q3h
2287	Q1w	Me	H	Q3i
2288	Q1w	Me	H	Q3j

2289	Q1w	Me	H	Q3k
2290	Q1w	Me	H	Q3l
2291	Q1w	Me	H	Q3m
2292	Q1w	Me	H	Q3n
2293	Q1w	Me	H	Q3o
2294	Q1w	Me	H	Q3p
2295	Q1w	Me	H	Q3q
2296	Q1w	Me	Me	Q3a
2297	Q1w	Me	Me	Q3b
2298	Q1w	Me	Me	Q3c
2299	Q1w	Me	Me	Q3d
2300	Q1w	Me	Me	Q3e
2301	Q1w	Me	Me	Q3f
2302	Q1w	Me	Me	Q3g
2303	Q1w	Me	Me	Q3h
2304	Q1w	Me	Me	Q3i
2305	Q1w	Me	Me	Q3j
2306	Q1w	Me	Me	Q3k
2307	Q1w	Me	Me	Q3l
2308	Q1w	Me	Me	Q3m
2309	Q1w	Me	Me	Q3n
2310	Q1w	Me	Me	Q3o
2311	Q1w	Me	Me	Q3p
2312	Q1w	Me	Me	Q3q
2313	Q1w	CF3	H	Q3a
2314	Q1w	CF3	H	Q3b
2315	Q1w	CF3	H	Q3c
2316	Q1w	CF3	H	Q3d
2317	Q1w	CF3	H	Q3e
2318	Q1w	CF3	H	Q3f
2319	Q1w	CF3	H	Q3g
2320	Q1w	CF3	H	Q3h
2321	Q1w	CF3	H	Q3i
2322	Q1w	CF3	H	Q3j
2323	Q1w	CF3	H	Q3k
2324	Q1w	CF3	H	Q3l

2325	Q1w	CF3	H	Q3m
2326	Q1w	CF3	H	Q3n
2327	Q1w	CF3	H	Q3o
2328	Q1w	CF3	H	Q3p
2329	Q1w	CF3	H	Q3q
2330	Q1w	CF3	Me	Q3a
2331	Q1w	CF3	Me	Q3b
2332	Q1w	CF3	Me	Q3c
2333	Q1w	CF3	Me	Q3d
2334	Q1w	CF3	Me	Q3e
2335	Q1w	CF3	Me	Q3f
2336	Q1w	CF3	Me	Q3g
2337	Q1w	CF3	Me	Q3h
2338	Q1w	CF3	Me	Q3i
2339	Q1w	CF3	Me	Q3j
2340	Q1w	CF3	Me	Q3k
2341	Q1w	CF3	Me	Q3l
2342	Q1w	CF3	Me	Q3m
2343	Q1w	CF3	Me	Q3n
2344	Q1w	CF3	Me	Q3o
2345	Q1w	CF3	Me	Q3p
2346	Q1w	CF3	Me	Q3q
2347	Q1x	H	H	Q3a
2348	Q1x	H	H	Q3b
2349	Q1x	H	H	Q3c
2350	Q1x	H	H	Q3d
2351	Q1x	H	H	Q3e
2352	Q1x	H	H	Q3f
2353	Q1x	H	H	Q3g
2354	Q1x	H	H	Q3h
2355	Q1x	H	H	Q3i
2356	Q1x	H	H	Q3j
2357	Q1x	H	H	Q3k
2358	Q1x	H	H	Q3l
2359	Q1x	H	H	Q3m
2360	Q1x	H	H	Q3n

2361	Q1x	H	H	Q3o
2362	Q1x	H	H	Q3p
2363	Q1x	H	H	Q3q
2364	Q1x	H	Me	Q3a
2365	Q1x	H	Me	Q3b
2366	Q1x	H	Me	Q3c
2367	Q1x	H	Me	Q3d
2368	Q1x	H	Me	Q3e
2369	Q1x	H	Me	Q3f
2370	Q1x	H	Me	Q3g
2371	Q1x	H	Me	Q3h
2372	Q1x	H	Me	Q3i
2373	Q1x	H	Me	Q3j
2374	Q1x	H	Me	Q3k
2375	Q1x	H	Me	Q3l
2376	Q1x	H	Me	Q3m
2377	Q1x	H	Me	Q3n
2378	Q1x	H	Me	Q3o
2379	Q1x	H	Me	Q3p
2380	Q1x	H	Me	Q3q
2381	Q1x	Me	H	Q3a
2382	Q1x	Me	H	Q3b
2383	Q1x	Me	H	Q3c
2384	Q1x	Me	H	Q3d
2385	Q1x	Me	H	Q3e
2386	Q1x	Me	H	Q3f
2387	Q1x	Me	H	Q3g
2388	Q1x	Me	H	Q3h
2389	Q1x	Me	H	Q3i
2390	Q1x	Me	H	Q3j
2391	Q1x	Me	H	Q3k
2392	Q1x	Me	H	Q3l
2393	Q1x	Me	H	Q3m
2394	Q1x	Me	H	Q3n
2395	Q1x	Me	H	Q3o
2396	Q1x	Me	H	Q3p

2397	Q1x	Me	H	Q3q
2398	Q1x	Me	Me	Q3a
2399	Q1x	Me	Me	Q3b
2400	Q1x	Me	Me	Q3c
2401	Q1x	Me	Me	Q3d
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2404	Q1x	Me	Me	Q3g
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2406	Q1x	Me	Me	Q3i
2407	Q1x	Me	Me	Q3j
2408	Q1x	Me	Me	Q3k
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2410	Q1x	Me	Me	Q3m
2411	Q1x	Me	Me	Q3n
2412	Q1x	Me	Me	Q3o
2413	Q1x	Me	Me	Q3p
2414	Q1x	Me	Me	Q3q
2415	Q1x	CF3	H	Q3a
2416	Q1x	CF3	H	Q3b
2417	Q1x	CF3	H	Q3c
2418	Q1x	CF3	H	Q3d
2419	Q1x	CF3	H	Q3e
2420	Q1x	CF3	H	Q3f
2421	Q1x	CF3	H	Q3g
2422	Q1x	CF3	H	Q3h
2423	Q1x	CF3	H	Q3i
2424	Q1x	CF3	H	Q3j
2425	Q1x	CF3	H	Q3k
2426	Q1x	CF3	H	Q3l
2427	Q1x	CF3	H	Q3m
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2431	Q1x	CF3	H	Q3q
2432	Q1x	CF3	Me	Q3a

2433	Q1x	CF3	Me	Q3b
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2436	Q1x	CF3	Me	Q3e
2437	Q1x	CF3	Me	Q3f
2438	Q1x	CF3	Me	Q3g
2439	Q1x	CF3	Me	Q3h
2440	Q1x	CF3	Me	Q3i
2441	Q1x	CF3	Me	Q3j
2442	Q1x	CF3	Me	Q3k
2443	Q1x	CF3	Me	Q3l
2444	Q1x	CF3	Me	Q3m
2445	Q1x	CF3	Me	Q3n
2446	Q1x	CF3	Me	Q3o
2447	Q1x	CF3	Me	Q3p
2448	Q1x	CF3	Me	Q3q
2449	Q1y	H	H	Q3a
2450	Q1y	H	H	Q3b
2451	Q1y	H	H	Q3c
2452	Q1y	H	H	Q3d
2453	Q1y	H	H	Q3e
2454	Q1y	H	H	Q3f
2455	Q1y	H	H	Q3g
2456	Q1y	H	H	Q3h
2457	Q1y	H	H	Q3i
2458	Q1y	H	H	Q3j
2459	Q1y	H	H	Q3k
2460	Q1y	H	H	Q3l
2461	Q1y	H	H	Q3m
2462	Q1y	H	H	Q3n
2463	Q1y	H	H	Q3o
2464	Q1y	H	H	Q3p
2465	Q1y	H	H	Q3q
2466	Q1y	H	Me	Q3a
2467	Q1y	H	Me	Q3b
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2469	Q1y	H	Me	Q3d
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2471	Q1y	H	Me	Q3f
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2485	Q1y	Me	H	Q3c
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2492	Q1y	Me	H	Q3j
2493	Q1y	Me	H	Q3k
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2500	Q1y	Me	Me	Q3a
2501	Q1y	Me	Me	Q3b
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2507	Q1y	Me	Me	Q3h
2508	Q1y	Me	Me	Q3i
2509	Q1y	Me	Me	Q3j
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2544	Q1y	CF3	Me	Q3k
2545	Q1y	CF3	Me	Q3l
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2549	Q1y	CF3	Me	Q3p
2550	Q1y	CF3	Me	Q3q
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2553	Q1z	H	H	Q3c
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2556	Q1z	H	H	Q3f
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2559	Q1z	H	H	Q3i
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2599	Q1z	Me	H	Q3o
2600	Q1z	Me	H	Q3p
2601	Q1z	Me	H	Q3q
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2618	Q1z	Me	Me	Q3q
2619	Q1z	CF3	H	Q3a
2620	Q1z	CF3	H	Q3b
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2627	Q1z	CF3	H	Q3i
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2635	Q1z	CF3	H	Q3q
2636	Q1z	CF3	Me	Q3a
2637	Q1z	CF3	Me	Q3b
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2647	Q1z	CF3	Me	Q3l
2648	Q1z	CF3	Me	Q3m

2649	Q1z	CF3	Me	Q3n
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2651	Q1z	CF3	Me	Q3p
2652	Q1z	CF3	Me	Q3q
2653	Q1a	H	H	Q3r
2654	Q1a	H	H	Q3s
2655	Q1a	H	H	Q3t
2656	Q1a	H	H	Q3u
2657	Q1a	H	Me	Q3r
2658	Q1a	H	Me	Q3s
2659	Q1a	H	Me	Q3t
2660	Q1a	H	Me	Q3u
2661	Q1a	Me	H	Q3r
2662	Q1a	Me	H	Q3s
2663	Q1a	Me	H	Q3t
2664	Q1a	Me	H	Q3u
2665	Q1a	Me	Me	Q3r
2666	Q1a	Me	Me	Q3s
2667	Q1a	Me	Me	Q3t
2668	Q1a	Me	Me	Q3u
2669	Q1a	CF3	H	Q3r
2670	Q1a	CF3	H	Q3s
2671	Q1a	CF3	H	Q3t
2672	Q1a	CF3	H	Q3u
2673	Q1a	CF3	Me	Q3r
2674	Q1a	CF3	Me	Q3s
2675	Q1a	CF3	Me	Q3t
2676	Q1a	CF3	Me	Q3u
2677	Q1b	H	H	Q3r
2678	Q1b	H	H	Q3s
2679	Q1b	H	H	Q3t
2680	Q1b	H	H	Q3u
2681	Q1b	H	Me	Q3r
2682	Q1b	H	Me	Q3s
2683	Q1b	H	Me	Q3t
2684	Q1b	H	Me	Q3u

2685	Q1b	Me	H	Q3r
2686	Q1b	Me	H	Q3s
2687	Q1b	Me	H	Q3t
2688	Q1b	Me	H	Q3u
2689	Q1b	Me	Me	Q3r
2690	Q1b	Me	Me	Q3s
2691	Q1b	Me	Me	Q3t
2692	Q1b	Me	Me	Q3u
2693	Q1b	CF3	H	Q3r
2694	Q1b	CF3	H	Q3s
2695	Q1b	CF3	H	Q3t
2696	Q1b	CF3	H	Q3u
2697	Q1b	CF3	Me	Q3r
2698	Q1b	CF3	Me	Q3s
2699	Q1b	CF3	Me	Q3t
2700	Q1b	CF3	Me	Q3u
2701	Q1c	H	H	Q3r
2702	Q1c	H	H	Q3s
2703	Q1c	H	H	Q3t
2704	Q1c	H	H	Q3u
2705	Q1c	H	Me	Q3r
2706	Q1c	H	Me	Q3s
2707	Q1c	H	Me	Q3t
2708	Q1c	H	Me	Q3u
2709	Q1c	Me	H	Q3r
2710	Q1c	Me	H	Q3s
2711	Q1c	Me	H	Q3t
2712	Q1c	Me	H	Q3u
2713	Q1c	Me	Me	Q3r
2714	Q1c	Me	Me	Q3s
2715	Q1c	Me	Me	Q3t
2716	Q1c	Me	Me	Q3u
2717	Q1c	CF3	H	Q3r
2718	Q1c	CF3	H	Q3s
2719	Q1c	CF3	H	Q3t
2720	Q1c	CF3	H	Q3u

2721	Q1c	CF3	Me	Q3r
2722	Q1c	CF3	Me	Q3s
2723	Q1c	CF3	Me	Q3t
2724	Q1c	CF3	Me	Q3u
2725	Q1d	H	H	Q3r
2726	Q1d	H	H	Q3s
2727	Q1d	H	H	Q3t
2728	Q1d	H	H	Q3u
2729	Q1d	H	Me	Q3r
2730	Q1d	H	Me	Q3s
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2737	Q1d	Me	Me	Q3r
2738	Q1d	Me	Me	Q3s
2739	Q1d	Me	Me	Q3t
2740	Q1d	Me	Me	Q3u
2741	Q1d	CF3	H	Q3r
2742	Q1d	CF3	H	Q3s
2743	Q1d	CF3	H	Q3t
2744	Q1d	CF3	H	Q3u
2745	Q1d	CF3	Me	Q3r
2746	Q1d	CF3	Me	Q3s
2747	Q1d	CF3	Me	Q3t
2748	Q1d	CF3	Me	Q3u
2749	Q1e	H	H	Q3r
2750	Q1e	H	H	Q3s
2751	Q1e	H	H	Q3t
2752	Q1e	H	H	Q3u
2753	Q1e	H	Me	Q3r
2754	Q1e	H	Me	Q3s
2755	Q1e	H	Me	Q3t
2756	Q1e	H	Me	Q3u

2757	Q1e	Me	H	Q3r
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2759	Q1e	Me	H	Q3t
2760	Q1e	Me	H	Q3u
2761	Q1e	Me	Me	Q3r
2762	Q1e	Me	Me	Q3s
2763	Q1e	Me	Me	Q3t
2764	Q1e	Me	Me	Q3u
2765	Q1e	CF3	H	Q3r
2766	Q1e	CF3	H	Q3s
2767	Q1e	CF3	H	Q3t
2768	Q1e	CF3	H	Q3u
2769	Q1e	CF3	Me	Q3r
2770	Q1e	CF3	Me	Q3s
2771	Q1e	CF3	Me	Q3t
2772	Q1e	CF3	Me	Q3u
2773	Q1f	H	H	Q3r
2774	Q1f	H	H	Q3s
2775	Q1f	H	H	Q3t
2776	Q1f	H	H	Q3u
2777	Q1f	H	Me	Q3r
2778	Q1f	H	Me	Q3s
2779	Q1f	H	Me	Q3t
2780	Q1f	H	Me	Q3u
2781	Q1f	Me	H	Q3r
2782	Q1f	Me	H	Q3s
2783	Q1f	Me	H	Q3t
2784	Q1f	Me	H	Q3u
2785	Q1f	Me	Me	Q3r
2786	Q1f	Me	Me	Q3s
2787	Q1f	Me	Me	Q3t
2788	Q1f	Me	Me	Q3u
2789	Q1f	CF3	H	Q3r
2790	Q1f	CF3	H	Q3s
2791	Q1f	CF3	H	Q3t
2792	Q1f	CF3	H	Q3u

2793	Q1f	CF3	Me	Q3r
2794	Q1f	CF3	Me	Q3s
2795	Q1f	CF3	Me	Q3t
2796	Q1f	CF3	Me	Q3u
2797	Q1g	H	H	Q3r
2798	Q1g	H	H	Q3s
2799	Q1g	H	H	Q3t
2800	Q1g	H	H	Q3u
2801	Q1g	H	Me	Q3r
2802	Q1g	H	Me	Q3s
2803	Q1g	H	Me	Q3t
2804	Q1g	H	Me	Q3u
2805	Q1g	Me	H	Q3r
2806	Q1g	Me	H	Q3s
2807	Q1g	Me	H	Q3t
2808	Q1g	Me	H	Q3u
2809	Q1g	Me	Me	Q3r
2810	Q1g	Me	Me	Q3s
2811	Q1g	Me	Me	Q3t
2812	Q1g	Me	Me	Q3u
2813	Q1g	CF3	H	Q3r
2814	Q1g	CF3	H	Q3s
2815	Q1g	CF3	H	Q3t
2816	Q1g	CF3	H	Q3u
2817	Q1g	CF3	Me	Q3r
2818	Q1g	CF3	Me	Q3s
2819	Q1g	CF3	Me	Q3t
2820	Q1g	CF3	Me	Q3u
2821	Q1h	H	H	Q3r
2822	Q1h	H	H	Q3s
2823	Q1h	H	H	Q3t
2824	Q1h	H	H	Q3u
2825	Q1h	H	Me	Q3r
2826	Q1h	H	Me	Q3s
2827	Q1h	H	Me	Q3t
2828	Q1h	H	Me	Q3u

2829	Q1h	Me	H	Q3r
2830	Q1h	Me	H	Q3s
2831	Q1h	Me	H	Q3t
2832	Q1h	Me	H	Q3u
2833	Q1h	Me	Me	Q3r
2834	Q1h	Me	Me	Q3s
2835	Q1h	Me	Me	Q3t
2836	Q1h	Me	Me	Q3u
2837	Q1h	CF3	H	Q3r
2838	Q1h	CF3	H	Q3s
2839	Q1h	CF3	H	Q3t
2840	Q1h	CF3	H	Q3u
2841	Q1h	CF3	Me	Q3r
2842	Q1h	CF3	Me	Q3s
2843	Q1h	CF3	Me	Q3t
2844	Q1h	CF3	Me	Q3u
2845	Q1i	H	H	Q3r
2846	Q1i	H	H	Q3s
2847	Q1i	H	H	Q3t
2848	Q1i	H	H	Q3u
2849	Q1i	H	Me	Q3r
2850	Q1i	H	Me	Q3s
2851	Q1i	H	Me	Q3t
2852	Q1i	H	Me	Q3u
2853	Q1i	Me	H	Q3r
2854	Q1i	Me	H	Q3s
2855	Q1i	Me	H	Q3t
2856	Q1i	Me	H	Q3u
2857	Q1i	Me	Me	Q3r
2858	Q1i	Me	Me	Q3s
2859	Q1i	Me	Me	Q3t
2860	Q1i	Me	Me	Q3u
2861	Q1i	CF3	H	Q3r
2862	Q1i	CF3	H	Q3s
2863	Q1i	CF3	H	Q3t
2864	Q1i	CF3	H	Q3u

2865	Q1i	CF3	Me	Q3r
2866	Q1i	CF3	Me	Q3s
2867	Q1i	CF3	Me	Q3t
2868	Q1i	CF3	Me	Q3u
2869	Q1j	H	H	Q3r
2870	Q1j	H	H	Q3s
2871	Q1j	H	H	Q3t
2872	Q1j	H	H	Q3u
2873	Q1j	H	Me	Q3r
2874	Q1j	H	Me	Q3s
2875	Q1j	H	Me	Q3t
2876	Q1j	H	Me	Q3u
2877	Q1j	Me	H	Q3r
2878	Q1j	Me	H	Q3s
2879	Q1j	Me	H	Q3t
2880	Q1j	Me	H	Q3u
2881	Q1j	Me	Me	Q3r
2882	Q1j	Me	Me	Q3s
2883	Q1j	Me	Me	Q3t
2884	Q1j	Me	Me	Q3u
2885	Q1j	CF3	H	Q3r
2886	Q1j	CF3	H	Q3s
2887	Q1j	CF3	H	Q3t
2888	Q1j	CF3	H	Q3u
2889	Q1j	CF3	Me	Q3r
2890	Q1j	CF3	Me	Q3s
2891	Q1j	CF3	Me	Q3t
2892	Q1j	CF3	Me	Q3u
2893	Q1k	H	H	Q3r
2894	Q1k	H	H	Q3s
2895	Q1k	H	H	Q3t
2896	Q1k	H	H	Q3u
2897	Q1k	H	Me	Q3r
2898	Q1k	H	Me	Q3s
2899	Q1k	H	Me	Q3t
2900	Q1k	H	Me	Q3u

2901	Q1k	Me	H	Q3r
2902	Q1k	Me	H	Q3s
2903	Q1k	Me	H	Q3t
2904	Q1k	Me	H	Q3u
2905	Q1k	Me	Me	Q3r
2906	Q1k	Me	Me	Q3s
2907	Q1k	Me	Me	Q3t
2908	Q1k	Me	Me	Q3u
2909	Q1k	CF3	H	Q3r
2910	Q1k	CF3	H	Q3s
2911	Q1k	CF3	H	Q3t
2912	Q1k	CF3	H	Q3u
2913	Q1k	CF3	Me	Q3r
2914	Q1k	CF3	Me	Q3s
2915	Q1k	CF3	Me	Q3t
2916	Q1k	CF3	Me	Q3u
2917	Q1l	H	H	Q3r
2918	Q1l	H	H	Q3s
2919	Q1l	H	H	Q3t
2920	Q1l	H	H	Q3u
2921	Q1l	H	Me	Q3r
2922	Q1l	H	Me	Q3s
2923	Q1l	H	Me	Q3t
2924	Q1l	H	Me	Q3u
2925	Q1l	Me	H	Q3r
2926	Q1l	Me	H	Q3s
2927	Q1l	Me	H	Q3t
2928	Q1l	Me	H	Q3u
2929	Q1l	Me	Me	Q3r
2930	Q1l	Me	Me	Q3s
2931	Q1l	Me	Me	Q3t
2932	Q1l	Me	Me	Q3u
2933	Q1l	CF3	H	Q3r
2934	Q1l	CF3	H	Q3s
2935	Q1l	CF3	H	Q3t
2936	Q1l	CF3	H	Q3u

2937	Q1l	CF3	Me	Q3r
2938	Q1l	CF3	Me	Q3s
2939	Q1l	CF3	Me	Q3t
2940	Q1l	CF3	Me	Q3u
2941	Q1m	H	H	Q3r
2942	Q1m	H	H	Q3s
2943	Q1m	H	H	Q3t
2944	Q1m	H	H	Q3u
2945	Q1m	H	Me	Q3r
2946	Q1m	H	Me	Q3s
2947	Q1m	H	Me	Q3t
2948	Q1m	H	Me	Q3u
2949	Q1m	Me	H	Q3r
2950	Q1m	Me	H	Q3s
2951	Q1m	Me	H	Q3t
2952	Q1m	Me	H	Q3u
2953	Q1m	Me	Me	Q3r
2954	Q1m	Me	Me	Q3s
2955	Q1m	Me	Me	Q3t
2956	Q1m	Me	Me	Q3u
2957	Q1m	CF3	H	Q3r
2958	Q1m	CF3	H	Q3s
2959	Q1m	CF3	H	Q3t
2960	Q1m	CF3	H	Q3u
2961	Q1m	CF3	Me	Q3r
2962	Q1m	CF3	Me	Q3s
2963	Q1m	CF3	Me	Q3t
2964	Q1m	CF3	Me	Q3u
2965	Q1n	H	H	Q3r
2966	Q1n	H	H	Q3s
2967	Q1n	H	H	Q3t
2968	Q1n	H	H	Q3u
2969	Q1n	H	Me	Q3r
2970	Q1n	H	Me	Q3s
2971	Q1n	H	Me	Q3t
2972	Q1n	H	Me	Q3u

2973	Q1n	Me	H	Q3r
2974	Q1n	Me	H	Q3s
2975	Q1n	Me	H	Q3t
2976	Q1n	Me	H	Q3u
2977	Q1n	Me	Me	Q3r
2978	Q1n	Me	Me	Q3s
2979	Q1n	Me	Me	Q3t
2980	Q1n	Me	Me	Q3u
2981	Q1n	CF3	H	Q3r
2982	Q1n	CF3	H	Q3s
2983	Q1n	CF3	H	Q3t
2984	Q1n	CF3	H	Q3u
2985	Q1n	CF3	Me	Q3r
2986	Q1n	CF3	Me	Q3s
2987	Q1n	CF3	Me	Q3t
2988	Q1n	CF3	Me	Q3u
2989	Q1o	H	H	Q3r
2990	Q1o	H	H	Q3s
2991	Q1o	H	H	Q3t
2992	Q1o	H	H	Q3u
2993	Q1o	H	Me	Q3r
2994	Q1o	H	Me	Q3s
2995	Q1o	H	Me	Q3t
2996	Q1o	H	Me	Q3u
2997	Q1o	Me	H	Q3r
2998	Q1o	Me	H	Q3s
2999	Q1o	Me	H	Q3t
3000	Q1o	Me	H	Q3u
3001	Q1o	Me	Me	Q3r
3002	Q1o	Me	Me	Q3s
3003	Q1o	Me	Me	Q3t
3004	Q1o	Me	Me	Q3u
3005	Q1o	CF3	H	Q3r
3006	Q1o	CF3	H	Q3s
3007	Q1o	CF3	H	Q3t
3008	Q1o	CF3	H	Q3u

3009	Q1o	CF3	Me	Q3r
3010	Q1o	CF3	Me	Q3s
3011	Q1o	CF3	Me	Q3t
3012	Q1o	CF3	Me	Q3u
3013	Q1p	H	H	Q3r
3014	Q1p	H	H	Q3s
3015	Q1p	H	H	Q3t
3016	Q1p	H	H	Q3u
3017	Q1p	H	Me	Q3r
3018	Q1p	H	Me	Q3s
3019	Q1p	H	Me	Q3t
3020	Q1p	H	Me	Q3u
3021	Q1p	Me	H	Q3r
3022	Q1p	Me	H	Q3s
3023	Q1p	Me	H	Q3t
3024	Q1p	Me	H	Q3u
3025	Q1p	Me	Me	Q3r
3026	Q1p	Me	Me	Q3s
3027	Q1p	Me	Me	Q3t
3028	Q1p	Me	Me	Q3u
3029	Q1p	CF3	H	Q3r
3030	Q1p	CF3	H	Q3s
3031	Q1p	CF3	H	Q3t
3032	Q1p	CF3	H	Q3u
3033	Q1p	CF3	Me	Q3r
3034	Q1p	CF3	Me	Q3s
3035	Q1p	CF3	Me	Q3t
3036	Q1p	CF3	Me	Q3u
3037	Q1q	H	H	Q3r
3038	Q1q	H	H	Q3s
3039	Q1q	H	H	Q3t
3040	Q1q	H	H	Q3u
3041	Q1q	H	Me	Q3r
3042	Q1q	H	Me	Q3s
3043	Q1q	H	Me	Q3t
3044	Q1q	H	Me	Q3u

3045	Q1q	Me	H	Q3r
3046	Q1q	Me	H	Q3s
3047	Q1q	Me	H	Q3t
3048	Q1q	Me	H	Q3u
3049	Q1q	Me	Me	Q3r
3050	Q1q	Me	Me	Q3s
3051	Q1q	Me	Me	Q3t
3052	Q1q	Me	Me	Q3u
3053	Q1q	CF3	H	Q3r
3054	Q1q	CF3	H	Q3s
3055	Q1q	CF3	H	Q3t
3056	Q1q	CF3	H	Q3u
3057	Q1q	CF3	Me	Q3r
3058	Q1q	CF3	Me	Q3s
3059	Q1q	CF3	Me	Q3t
3060	Q1q	CF3	Me	Q3u
3061	Q1r	H	H	Q3r
3062	Q1r	H	H	Q3s
3063	Q1r	H	H	Q3t
3064	Q1r	H	H	Q3u
3065	Q1r	H	Me	Q3r
3066	Q1r	H	Me	Q3s
3067	Q1r	H	Me	Q3t
3068	Q1r	H	Me	Q3u
3069	Q1r	Me	H	Q3r
3070	Q1r	Me	H	Q3s
3071	Q1r	Me	H	Q3t
3072	Q1r	Me	H	Q3u
3073	Q1r	Me	Me	Q3r
3074	Q1r	Me	Me	Q3s
3075	Q1r	Me	Me	Q3t
3076	Q1r	Me	Me	Q3u
3077	Q1r	CF3	H	Q3r
3078	Q1r	CF3	H	Q3s
3079	Q1r	CF3	H	Q3t
3080	Q1r	CF3	H	Q3u

3081	Q1r	CF3	Me	Q3r
3082	Q1r	CF3	Me	Q3s
3083	Q1r	CF3	Me	Q3t
3084	Q1r	CF3	Me	Q3u
3085	Q1s	H	H	Q3r
3086	Q1s	H	H	Q3s
3087	Q1s	H	H	Q3t
3088	Q1s	H	H	Q3u
3089	Q1s	H	Me	Q3r
3090	Q1s	H	Me	Q3s
3091	Q1s	H	Me	Q3t
3092	Q1s	H	Me	Q3u
3093	Q1s	Me	H	Q3r
3094	Q1s	Me	H	Q3s
3095	Q1s	Me	H	Q3t
3096	Q1s	Me	H	Q3u
3097	Q1s	Me	Me	Q3r
3098	Q1s	Me	Me	Q3s
3099	Q1s	Me	Me	Q3t
3100	Q1s	Me	Me	Q3u
3101	Q1s	CF3	H	Q3r
3102	Q1s	CF3	H	Q3s
3103	Q1s	CF3	H	Q3t
3104	Q1s	CF3	H	Q3u
3105	Q1s	CF3	Me	Q3r
3106	Q1s	CF3	Me	Q3s
3107	Q1s	CF3	Me	Q3t
3108	Q1s	CF3	Me	Q3u
3109	Q1t	H	H	Q3r
3110	Q1t	H	H	Q3s
3111	Q1t	H	H	Q3t
3112	Q1t	H	H	Q3u
3113	Q1t	H	Me	Q3r
3114	Q1t	H	Me	Q3s
3115	Q1t	H	Me	Q3t
3116	Q1t	H	Me	Q3u

3117	Q1t	Me	H	Q3r
3118	Q1t	Me	H	Q3s
3119	Q1t	Me	H	Q3t
3120	Q1t	Me	H	Q3u
3121	Q1t	Me	Me	Q3r
3122	Q1t	Me	Me	Q3s
3123	Q1t	Me	Me	Q3t
3124	Q1t	Me	Me	Q3u
3125	Q1t	CF3	H	Q3r
3126	Q1t	CF3	H	Q3s
3127	Q1t	CF3	H	Q3t
3128	Q1t	CF3	H	Q3u
3129	Q1t	CF3	Me	Q3r
3130	Q1t	CF3	Me	Q3s
3131	Q1t	CF3	Me	Q3t
3132	Q1t	CF3	Me	Q3u
3133	Q1u	H	H	Q3r
3134	Q1u	H	H	Q3s
3135	Q1u	H	H	Q3t
3136	Q1u	H	H	Q3u
3137	Q1u	H	Me	Q3r
3138	Q1u	H	Me	Q3s
3139	Q1u	H	Me	Q3t
3140	Q1u	H	Me	Q3u
3141	Q1u	Me	H	Q3r
3142	Q1u	Me	H	Q3s
3143	Q1u	Me	H	Q3t
3144	Q1u	Me	H	Q3u
3145	Q1u	Me	Me	Q3r
3146	Q1u	Me	Me	Q3s
3147	Q1u	Me	Me	Q3t
3148	Q1u	Me	Me	Q3u
3149	Q1u	CF3	H	Q3r
3150	Q1u	CF3	H	Q3s
3151	Q1u	CF3	H	Q3t
3152	Q1u	CF3	H	Q3u

3153	Q1u	CF3	Me	Q3r
3154	Q1u	CF3	Me	Q3s
3155	Q1u	CF3	Me	Q3t
3156	Q1u	CF3	Me	Q3u
3157	Q1v	H	H	Q3r
3158	Q1v	H	H	Q3s
3159	Q1v	H	H	Q3t
3160	Q1v	H	H	Q3u
3161	Q1v	H	Me	Q3r
3162	Q1v	H	Me	Q3s
3163	Q1v	H	Me	Q3t
3164	Q1v	H	Me	Q3u
3165	Q1v	Me	H	Q3r
3166	Q1v	Me	H	Q3s
3167	Q1v	Me	H	Q3t
3168	Q1v	Me	H	Q3u
3169	Q1v	Me	Me	Q3r
3170	Q1v	Me	Me	Q3s
3171	Q1v	Me	Me	Q3t
3172	Q1v	Me	Me	Q3u
3173	Q1v	CF3	H	Q3r
3174	Q1v	CF3	H	Q3s
3175	Q1v	CF3	H	Q3t
3176	Q1v	CF3	H	Q3u
3177	Q1v	CF3	Me	Q3r
3178	Q1v	CF3	Me	Q3s
3179	Q1v	CF3	Me	Q3t
3180	Q1v	CF3	Me	Q3u
3181	Q1w	H	H	Q3r
3182	Q1w	H	H	Q3s
3183	Q1w	H	H	Q3t
3184	Q1w	H	H	Q3u
3185	Q1w	H	Me	Q3r
3186	Q1w	H	Me	Q3s
3187	Q1w	H	Me	Q3t
3188	Q1w	H	Me	Q3u

3189	Q1w	Me	H	Q3r
3190	Q1w	Me	H	Q3s
3191	Q1w	Me	H	Q3t
3192	Q1w	Me	H	Q3u
3193	Q1w	Me	Me	Q3r
3194	Q1w	Me	Me	Q3s
3195	Q1w	Me	Me	Q3t
3196	Q1w	Me	Me	Q3u
3197	Q1w	CF3	H	Q3r
3198	Q1w	CF3	H	Q3s
3199	Q1w	CF3	H	Q3t
3200	Q1w	CF3	H	Q3u
3201	Q1w	CF3	Me	Q3r
3202	Q1w	CF3	Me	Q3s
3203	Q1w	CF3	Me	Q3t
3204	Q1w	CF3	Me	Q3u
3205	Q1x	H	H	Q3r
3206	Q1x	H	H	Q3s
3207	Q1x	H	H	Q3t
3208	Q1x	H	H	Q3u
3209	Q1x	H	Me	Q3r
3210	Q1x	H	Me	Q3s
3211	Q1x	H	Me	Q3t
3212	Q1x	H	Me	Q3u
3213	Q1x	Me	H	Q3r
3214	Q1x	Me	H	Q3s
3215	Q1x	Me	H	Q3t
3216	Q1x	Me	H	Q3u
3217	Q1x	Me	Me	Q3r
3218	Q1x	Me	Me	Q3s
3219	Q1x	Me	Me	Q3t
3220	Q1x	Me	Me	Q3u
3221	Q1x	CF3	H	Q3r
3222	Q1x	CF3	H	Q3s
3223	Q1x	CF3	H	Q3t
3224	Q1x	CF3	H	Q3u

3225	Q1x	CF3	Me	Q3r
3226	Q1x	CF3	Me	Q3s
3227	Q1x	CF3	Me	Q3t
3228	Q1x	CF3	Me	Q3u
3229	Q1y	H	H	Q3r
3230	Q1y	H	H	Q3s
3231	Q1y	H	H	Q3t
3232	Q1y	H	H	Q3u
3233	Q1y	H	Me	Q3r
3234	Q1y	H	Me	Q3s
3235	Q1y	H	Me	Q3t
3236	Q1y	H	Me	Q3u
3237	Q1y	Me	H	Q3r
3238	Q1y	Me	H	Q3s
3239	Q1y	Me	H	Q3t
3240	Q1y	Me	H	Q3u
3241	Q1y	Me	Me	Q3r
3242	Q1y	Me	Me	Q3s
3243	Q1y	Me	Me	Q3t
3244	Q1y	Me	Me	Q3u
3245	Q1y	CF3	H	Q3r
3246	Q1y	CF3	H	Q3s
3247	Q1y	CF3	H	Q3t
3248	Q1y	CF3	H	Q3u
3249	Q1y	CF3	Me	Q3r
3250	Q1y	CF3	Me	Q3s
3251	Q1y	CF3	Me	Q3t
3252	Q1y	CF3	Me	Q3u
3253	Q1z	H	H	Q3r
3254	Q1z	H	H	Q3s
3255	Q1z	H	H	Q3t
3256	Q1z	H	H	Q3u
3257	Q1z	H	Me	Q3r
3258	Q1z	H	Me	Q3s
3259	Q1z	H	Me	Q3t
3260	Q1z	H	Me	Q3u

3261	Q1z	Me	H	Q3r
3262	Q1z	Me	H	Q3s
3263	Q1z	Me	H	Q3t
3264	Q1z	Me	H	Q3u
3265	Q1z	Me	Me	Q3r
3266	Q1z	Me	Me	Q3s
3267	Q1z	Me	Me	Q3t
3268	Q1z	Me	Me	Q3u
3269	Q1z	CF3	H	Q3r
3270	Q1z	CF3	H	Q3s
3271	Q1z	CF3	H	Q3t
3272	Q1z	CF3	H	Q3u
3273	Q1z	CF3	Me	Q3r
3274	Q1z	CF3	Me	Q3s
3275	Q1z	CF3	Me	Q3t
3276	Q1z	CF3	Me	Q3u

131) The compounds wherein R^1 , R^2 , R^3 and R^4 are any of the following combinations in Table 2, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof. The symbols in Table 2
5 denote the following substituents.

[Ka 11]

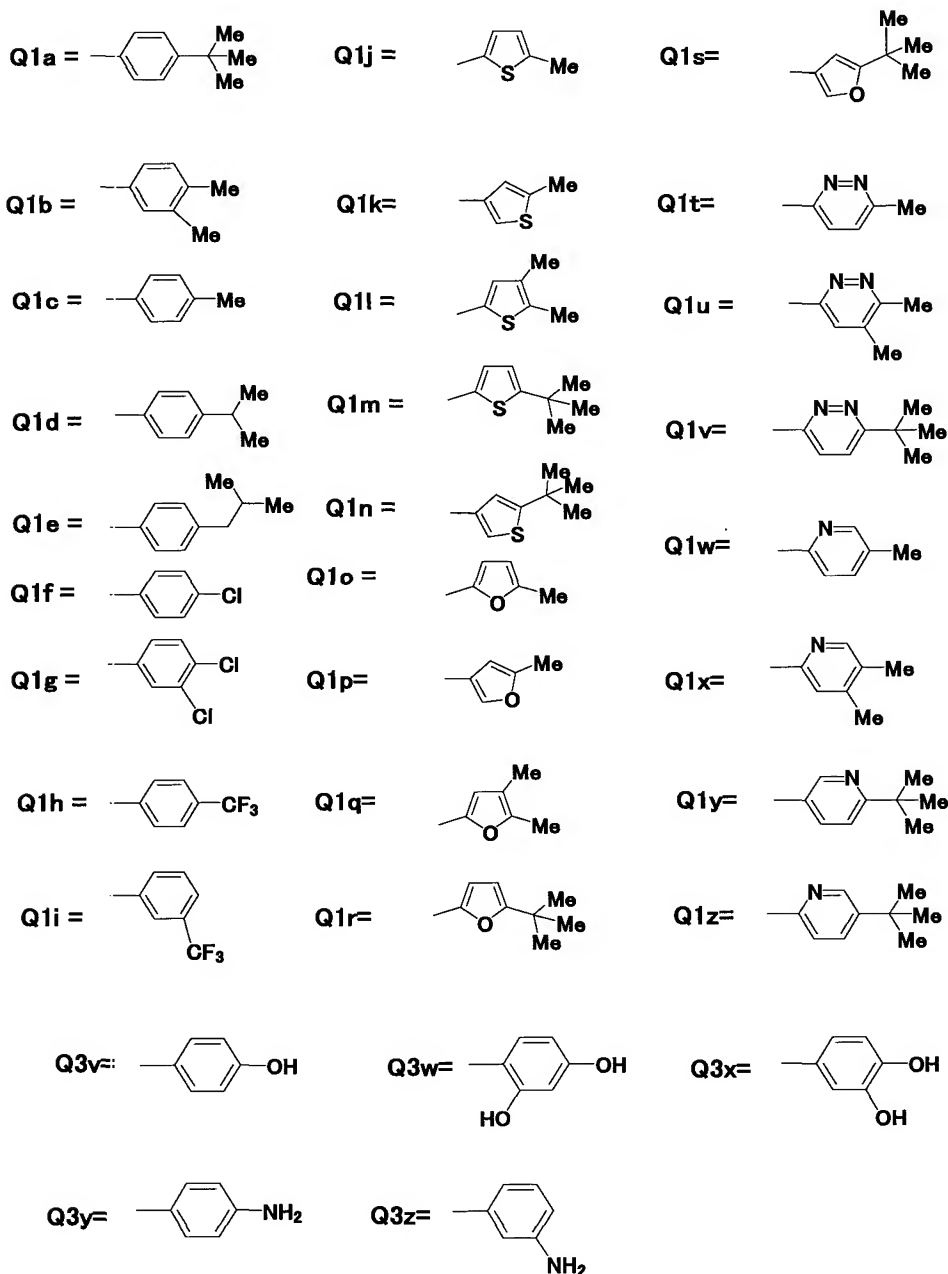


Table 2

No	R ¹	R ²	R ³	R ⁴
1	Q1a	H	H	Q3v
2	Q1a	H	H	Q3w
3	Q1a	H	H	Q3x
4	Q1a	H	H	Q3y
5	Q1a	H	H	Q3z
6	Q1a	H	Me	Q3v
7	Q1a	H	Me	Q3w
8	Q1a	H	Me	Q3x
9	Q1a	H	Me	Q3y
10	Q1a	H	Me	Q3z
11	Q1a	Me	H	Q3v
12	Q1a	Me	H	Q3w
13	Q1a	Me	H	Q3x
14	Q1a	Me	H	Q3y
15	Q1a	Me	H	Q3z
16	Q1a	Me	Me	Q3v
17	Q1a	Me	Me	Q3w
18	Q1a	Me	Me	Q3x
19	Q1a	Me	Me	Q3y
20	Q1a	Me	Me	Q3z
21	Q1a	CF3	H	Q3v
22	Q1a	CF3	H	Q3w
23	Q1a	CF3	H	Q3x
24	Q1a	CF3	H	Q3y
25	Q1a	CF3	H	Q3z
26	Q1a	CF3	Me	Q3v
27	Q1a	CF3	Me	Q3w
28	Q1a	CF3	Me	Q3x
29	Q1a	CF3	Me	Q3y
30	Q1a	CF3	Me	Q3z
31	Q1b	H	H	Q3v
32	Q1b	H	H	Q3w
33	Q1b	H	H	Q3x

34	Q1b	H	H	Q3y
35	Q1b	H	H	Q3z
36	Q1b	H	Me	Q3v
37	Q1b	H	Me	Q3w
38	Q1b	H	Me	Q3x
39	Q1b	H	Me	Q3y
40	Q1b	H	Me	Q3z
41	Q1b	Me	H	Q3v
42	Q1b	Me	H	Q3w
43	Q1b	Me	H	Q3x
44	Q1b	Me	H	Q3y
45	Q1b	Me	H	Q3z
46	Q1b	Me	Me	Q3v
47	Q1b	Me	Me	Q3w
48	Q1b	Me	Me	Q3x
49	Q1b	Me	Me	Q3y
50	Q1b	Me	Me	Q3z
51	Q1b	CF3	H	Q3v
52	Q1b	CF3	H	Q3w
53	Q1b	CF3	H	Q3x
54	Q1b	CF3	H	Q3y
55	Q1b	CF3	H	Q3z
56	Q1b	CF3	Me	Q3v
57	Q1b	CF3	Me	Q3w
58	Q1b	CF3	Me	Q3x
59	Q1b	CF3	Me	Q3y
60	Q1b	CF3	Me	Q3z
61	Q1c	H	H	Q3v
62	Q1c	H	H	Q3w
63	Q1c	H	H	Q3x
64	Q1c	H	H	Q3y
65	Q1c	H	H	Q3z
66	Q1c	H	Me	Q3v
67	Q1c	H	Me	Q3w
68	Q1c	H	Me	Q3x
69	Q1c	H	Me	Q3y

70	Q1c	H	Me	Q3z
71	Q1c	Me	H	Q3v
72	Q1c	Me	H	Q3w
73	Q1c	Me	H	Q3x
74	Q1c	Me	H	Q3y
75	Q1c	Me	H	Q3z
76	Q1c	Me	Me	Q3v
77	Q1c	Me	Me	Q3w
78	Q1c	Me	Me	Q3x
79	Q1c	Me	Me	Q3y
80	Q1c	Me	Me	Q3z
81	Q1c	CF3	H	Q3v
82	Q1c	CF3	H	Q3w
83	Q1c	CF3	H	Q3x
84	Q1c	CF3	H	Q3y
85	Q1c	CF3	H	Q3z
86	Q1c	CF3	Me	Q3v
87	Q1c	CF3	Me	Q3w
88	Q1c	CF3	Me	Q3x
89	Q1c	CF3	Me	Q3y
90	Q1c	CF3	Me	Q3z
91	Q1d	H	H	Q3v
92	Q1d	H	H	Q3w
93	Q1d	H	H	Q3x
94	Q1d	H	H	Q3y
95	Q1d	H	H	Q3z
96	Q1d	H	Me	Q3v
97	Q1d	H	Me	Q3w
98	Q1d	H	Me	Q3x
99	Q1d	H	Me	Q3y
100	Q1d	H	Me	Q3z
101	Q1d	Me	H	Q3v
102	Q1d	Me	H	Q3w
103	Q1d	Me	H	Q3x
104	Q1d	Me	H	Q3y
105	Q1d	Me	H	Q3z

106	Q1d	Me	Me	Q3v
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108	Q1d	Me	Me	Q3x
109	Q1d	Me	Me	Q3y
110	Q1d	Me	Me	Q3z
111	Q1d	CF3	H	Q3v
112	Q1d	CF3	H	Q3w
113	Q1d	CF3	H	Q3x
114	Q1d	CF3	H	Q3y
115	Q1d	CF3	H	Q3z
116	Q1d	CF3	Me	Q3v
117	Q1d	CF3	Me	Q3w
118	Q1d	CF3	Me	Q3x
119	Q1d	CF3	Me	Q3y
120	Q1d	CF3	Me	Q3z
121	Q1e	H	H	Q3v
122	Q1e	H	H	Q3w
123	Q1e	H	H	Q3x
124	Q1e	H	H	Q3y
125	Q1e	H	H	Q3z
126	Q1e	H	Me	Q3v
127	Q1e	H	Me	Q3w
128	Q1e	H	Me	Q3x
129	Q1e	H	Me	Q3y
130	Q1e	H	Me	Q3z
131	Q1e	Me	H	Q3v
132	Q1e	Me	H	Q3w
133	Q1e	Me	H	Q3x
134	Q1e	Me	H	Q3y
135	Q1e	Me	H	Q3z
136	Q1e	Me	Me	Q3v
137	Q1e	Me	Me	Q3w
138	Q1e	Me	Me	Q3x
139	Q1e	Me	Me	Q3y
140	Q1e	Me	Me	Q3z
141	Q1e	CF3	H	Q3v

142	Q1e	CF3	H	Q3w
143	Q1e	CF3	H	Q3x
144	Q1e	CF3	H	Q3y
145	Q1e	CF3	H	Q3z
146	Q1e	CF3	Me	Q3v
147	Q1e	CF3	Me	Q3w
148	Q1e	CF3	Me	Q3x
149	Q1e	CF3	Me	Q3y
150	Q1e	CF3	Me	Q3z
151	Q1f	H	H	Q3v
152	Q1f	H	H	Q3w
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154	Q1f	H	H	Q3y
155	Q1f	H	H	Q3z
156	Q1f	H	Me	Q3v
157	Q1f	H	Me	Q3w
158	Q1f	H	Me	Q3x
159	Q1f	H	Me	Q3y
160	Q1f	H	Me	Q3z
161	Q1f	Me	H	Q3v
162	Q1f	Me	H	Q3w
163	Q1f	Me	H	Q3x
164	Q1f	Me	H	Q3y
165	Q1f	Me	H	Q3z
166	Q1f	Me	Me	Q3v
167	Q1f	Me	Me	Q3w
168	Q1f	Me	Me	Q3x
169	Q1f	Me	Me	Q3y
170	Q1f	Me	Me	Q3z
171	Q1f	CF3	H	Q3v
172	Q1f	CF3	H	Q3w
173	Q1f	CF3	H	Q3x
174	Q1f	CF3	H	Q3y
175	Q1f	CF3	H	Q3z
176	Q1f	CF3	Me	Q3v
177	Q1f	CF3	Me	Q3w

178	Q1f	CF3	Me	Q3x
179	Q1f	CF3	Me	Q3y
180	Q1f	CF3	Me	Q3z
181	Q1g	H	H	Q3v
182	Q1g	H	H	Q3w
183	Q1g	H	H	Q3x
184	Q1g	H	H	Q3y
185	Q1g	H	H	Q3z
186	Q1g	H	Me	Q3v
187	Q1g	H	Me	Q3w
188	Q1g	H	Me	Q3x
189	Q1g	H	Me	Q3y
190	Q1g	H	Me	Q3z
191	Q1g	Me	H	Q3v
192	Q1g	Me	H	Q3w
193	Q1g	Me	H	Q3x
194	Q1g	Me	H	Q3y
195	Q1g	Me	H	Q3z
196	Q1g	Me	Me	Q3v
197	Q1g	Me	Me	Q3w
198	Q1g	Me	Me	Q3x
199	Q1g	Me	Me	Q3y
200	Q1g	Me	Me	Q3z
201	Q1g	CF3	H	Q3v
202	Q1g	CF3	H	Q3w
203	Q1g	CF3	H	Q3x
204	Q1g	CF3	H	Q3y
205	Q1g	CF3	H	Q3z
206	Q1g	CF3	Me	Q3v
207	Q1g	CF3	Me	Q3w
208	Q1g	CF3	Me	Q3x
209	Q1g	CF3	Me	Q3y
210	Q1g	CF3	Me	Q3z
211	Q1h	H	H	Q3v
212	Q1h	H	H	Q3w
213	Q1h	H	H	Q3x

214	Q1h	H	H	Q3y
215	Q1h	H	H	Q3z
216	Q1h	H	Me	Q3v
217	Q1h	H	Me	Q3w
218	Q1h	H	Me	Q3x
219	Q1h	H	Me	Q3y
220	Q1h	H	Me	Q3z
221	Q1h	Me	H	Q3v
222	Q1h	Me	H	Q3w
223	Q1h	Me	H	Q3x
224	Q1h	Me	H	Q3y
225	Q1h	Me	H	Q3z
226	Q1h	Me	Me	Q3v
227	Q1h	Me	Me	Q3w
228	Q1h	Me	Me	Q3x
229	Q1h	Me	Me	Q3y
230	Q1h	Me	Me	Q3z
231	Q1h	CF3	H	Q3v
232	Q1h	CF3	H	Q3w
233	Q1h	CF3	H	Q3x
234	Q1h	CF3	H	Q3y
235	Q1h	CF3	H	Q3z
236	Q1h	CF3	Me	Q3v
237	Q1h	CF3	Me	Q3w
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239	Q1h	CF3	Me	Q3y
240	Q1h	CF3	Me	Q3z
241	Q1i	H	H	Q3v
242	Q1i	H	H	Q3w
243	Q1i	H	H	Q3x
244	Q1i	H	H	Q3y
245	Q1i	H	H	Q3z
246	Q1i	H	Me	Q3v
247	Q1i	H	Me	Q3w
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249	Q1i	H	Me	Q3y

250	Q1i	H	Me	Q3z
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252	Q1i	Me	H	Q3w
253	Q1i	Me	H	Q3x
254	Q1i	Me	H	Q3y
255	Q1i	Me	H	Q3z
256	Q1i	Me	Me	Q3v
257	Q1i	Me	Me	Q3w
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259	Q1i	Me	Me	Q3y
260	Q1i	Me	Me	Q3z
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262	Q1i	CF3	H	Q3w
263	Q1i	CF3	H	Q3x
264	Q1i	CF3	H	Q3y
265	Q1i	CF3	H	Q3z
266	Q1I	CF3	Me	Q3v
267	Q1I	CF3	Me	Q3w
268	Q1I	CF3	Me	Q3X
269	Q1I	CF3	Me	Q3y
270	Q1I	CF3	Me	Q3z
271	Q1j	H	H	Q3v
272	Q1j	H	H	Q3w
273	Q1j	H	H	Q3X
274	Q1j	H	H	Q3y
275	Q1j	H	H	Q3z
276	Q1j	H	Me	Q3v
277	Q1j	H	Me	Q3w
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279	Q1j	H	Me	Q3y
280	Q1j	H	Me	Q3z
281	Q1j	Me	H	Q3v
282	Q1j	Me	H	Q3w
283	Q1j	Me	H	Q3x
284	Q1j	Me	H	Q3y
285	Q1j	Me	H	Q3z

286	Q1j	Me	Me	Q3v
287	Q1j	Me	Me	Q3w
288	Q1j	Me	Me	Q3x
289	Q1j	Me	Me	Q3y
290	Q1j	Me	Me	Q3z
291	Q1j	CF3	H	Q3v
292	Q1j	CF3	H	Q3w
293	Q1j	CF3	H	Q3x
294	Q1j	CF3	H	Q3y
295	Q1j	CF3	H	Q3z
296	Q1j	CF3	Me	Q3v
297	Q1j	CF3	Me	Q3w
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299	Q1j	CF3	Me	Q3y
300	Q1j	CF3	Me	Q3z
301	Q1k	H	H	Q3v
302	Q1k	H	H	Q3w
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304	Q1k	H	H	Q3y
305	Q1k	H	H	Q3z
306	Q1k	H	Me	Q3v
307	Q1k	H	Me	Q3w
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312	Q1k	Me	H	Q3w
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314	Q1k	Me	H	Q3y
315	Q1k	Me	H	Q3z
316	Q1k	Me	Me	Q3v
317	Q1k	Me	Me	Q3w
318	Q1k	Me	Me	Q3x
319	Q1k	Me	Me	Q3y
320	Q1k	Me	Me	Q3z
321	Q1k	CF3	H	Q3v

322	Q1k	CF3	H	Q3w
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325	Q1k	CF3	H	Q3z
326	Q1k	CF3	Me	Q3v
327	Q1k	CF3	Me	Q3w
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329	Q1k	CF3	Me	Q3y
330	Q1k	CF3	Me	Q3z
331	Q1l	H	H	Q3v
332	Q1l	H	H	Q3w
333	Q1l	H	H	Q3x
334	Q1l	H	H	Q3y
335	Q1l	H	H	Q3z
336	Q1l	H	Me	Q3v
337	Q1l	H	Me	Q3w
338	Q1l	H	Me	Q3x
339	Q1l	H	Me	Q3y
340	Q1l	H	Me	Q3z
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342	Q1l	Me	H	Q3w
343	Q1l	Me	H	Q3x
344	Q1l	Me	H	Q3y
345	Q1l	Me	H	Q3z
346	Q1l	Me	Me	Q3v
347	Q1l	Me	Me	Q3w
348	Q1l	Me	Me	Q3x
349	Q1l	Me	Me	Q3y
350	Q1l	Me	Me	Q3z
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353	Q1l	CF3	H	Q3x
354	Q1l	CF3	H	Q3y
355	Q1l	CF3	H	Q3z
356	Q1l	CF3	Me	Q3v
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358	Q1l	CF3	Me	Q3x
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360	Q1l	CF3	Me	Q3z
361	Q1m	H	H	Q3v
362	Q1m	H	H	Q3w
363	Q1m	H	H	Q3x
364	Q1m	H	H	Q3y
365	Q1m	H	H	Q3z
366	Q1m	H	Me	Q3v
367	Q1m	H	Me	Q3w
368	Q1m	H	Me	Q3x
369	Q1m	H	Me	Q3y
370	Q1m	H	Me	Q3z
371	Q1m	Me	H	Q3v
372	Q1m	Me	H	Q3w
373	Q1m	Me	H	Q3x
374	Q1m	Me	H	Q3y
375	Q1m	Me	H	Q3z
376	Q1m	Me	Me	Q3v
377	Q1m	Me	Me	Q3w
378	Q1m	Me	Me	Q3x
379	Q1m	Me	Me	Q3y
380	Q1m	Me	Me	Q3z
381	Q1m	CF3	H	Q3v
382	Q1m	CF3	H	Q3w
383	Q1m	CF3	H	Q3x
384	Q1m	CF3	H	Q3y
385	Q1m	CF3	H	Q3z
386	Q1m	CF3	Me	Q3v
387	Q1m	CF3	Me	Q3w
388	Q1m	CF3	Me	Q3x
389	Q1m	CF3	Me	Q3y
390	Q1m	CF3	Me	Q3z
391	Q1n	H	H	Q3v
392	Q1n	H	H	Q3w
393	Q1n	H	H	Q3x

394	Q1n	H	H	Q3y
395	Q1n	H	H	Q3z
396	Q1n	H	Me	Q3v
397	Q1n	H	Me	Q3w
398	Q1n	H	Me	Q3x
399	Q1n	H	Me	Q3y
400	Q1n	H	Me	Q3z
401	Q1n	Me	H	Q3v
402	Q1n	Me	H	Q3w
403	Q1n	Me	H	Q3x
404	Q1n	Me	H	Q3y
405	Q1n	Me	H	Q3z
406	Q1n	Me	Me	Q3v
407	Q1n	Me	Me	Q3w
408	Q1n	Me	Me	Q3x
409	Q1n	Me	Me	Q3y
410	Q1n	Me	Me	Q3z
411	Q1n	CF3	H	Q3v
412	Q1n	CF3	H	Q3w
413	Q1n	CF3	H	Q3x
414	Q1n	CF3	H	Q3y
415	Q1n	CF3	H	Q3z
416	Q1n	CF3	Me	Q3v
417	Q1n	CF3	Me	Q3w
418	Q1n	CF3	Me	Q3x
419	Q1n	CF3	Me	Q3y
420	Q1n	CF3	Me	Q3z
421	Q1o	H	H	Q3v
422	Q1o	H	H	Q3w
423	Q1o	H	H	Q3x
424	Q1o	H	H	Q3y
425	Q1o	H	H	Q3z
426	Q1o	H	Me	Q3v
427	Q1o	H	Me	Q3w
428	Q1o	H	Me	Q3x
429	Q1o	H	Me	Q3y

430	Q1o	H	Me	Q3z
431	Q1o	Me	H	Q3v
432	Q1o	Me	H	Q3w
433	Q1o	Me	H	Q3x
434	Q1o	Me	H	Q3y
435	Q1o	Me	H	Q3z
436	Q1o	Me	Me	Q3v
437	Q1o	Me	Me	Q3w
438	Q1o	Me	Me	Q3x
439	Q1o	Me	Me	Q3y
440	Q1o	Me	Me	Q3z
441	Q1o	CF3	H	Q3v
442	Q1o	CF3	H	Q3w
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444	Q1o	CF3	H	Q3y
445	Q1o	CF3	H	Q3z
446	Q1o	CF3	Me	Q3v
447	Q1o	CF3	Me	Q3w
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449	Q1o	CF3	Me	Q3y
450	Q1o	CF3	Me	Q3z
451	Q1p	H	H	Q3v
452	Q1p	H	H	Q3w
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455	Q1p	H	H	Q3z
456	Q1p	H	Me	Q3v
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459	Q1p	H	Me	Q3y
460	Q1p	H	Me	Q3z
461	Q1p	Me	H	Q3v
462	Q1p	Me	H	Q3w
463	Q1p	Me	H	Q3x
464	Q1p	Me	H	Q3y
465	Q1p	Me	H	Q3z

466	Q1p	Me	Me	Q3v
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468	Q1p	Me	Me	Q3x
469	Q1p	Me	Me	Q3y
470	Q1p	Me	Me	Q3z
471	Q1p	CF3	H	Q3v
472	Q1p	CF3	H	Q3w
473	Q1p	CF3	H	Q3x
474	Q1p	CF3	H	Q3y
475	Q1p	CF3	H	Q3z
476	Q1p	CF3	Me	Q3v
477	Q1p	CF3	Me	Q3w
478	Q1p	CF3	Me	Q3x
479	Q1p	CF3	Me	Q3y
480	Q1p	CF3	Me	Q3z
481	Q1q	H	H	Q3v
482	Q1q	H	H	Q3w
483	Q1q	H	H	Q3x
484	Q1q	H	H	Q3y
485	Q1q	H	H	Q3z
486	Q1q	H	Me	Q3v
487	Q1q	H	Me	Q3w
488	Q1q	H	Me	Q3x
489	Q1q	H	Me	Q3y
490	Q1q	H	Me	Q3z
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492	Q1q	Me	H	Q3w
493	Q1q	Me	H	Q3x
494	Q1q	Me	H	Q3y
495	Q1q	Me	H	Q3z
496	Q1q	Me	Me	Q3v
497	Q1q	Me	Me	Q3w
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499	Q1q	Me	Me	Q3y
500	Q1q	Me	Me	Q3z
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502	Q1q	CF3	H	Q3w
503	Q1q	CF3	H	Q3x
504	Q1q	CF3	H	Q3y
505	Q1q	CF3	H	Q3z
506	Q1q	CF3	Me	Q3v
507	Q1q	CF3	Me	Q3w
508	Q1q	CF3	Me	Q3x
509	Q1q	CF3	Me	Q3y
510	Q1q	CF3	Me	Q3z
511	Q1r	H	H	Q3v
512	Q1r	H	H	Q3w
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514	Q1r	H	H	Q3y
515	Q1r	H	H	Q3z
516	Q1r	H	Me	Q3v
517	Q1r	H	Me	Q3w
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519	Q1r	H	Me	Q3y
520	Q1r	H	Me	Q3z
521	Q1r	Me	H	Q3v
522	Q1r	Me	H	Q3w
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524	Q1r	Me	H	Q3y
525	Q1r	Me	H	Q3z
526	Q1r	Me	Me	Q3v
527	Q1r	Me	Me	Q3w
528	Q1r	Me	Me	Q3x
529	Q1r	Me	Me	Q3y
530	Q1r	Me	Me	Q3z
531	Q1r	CF3	H	Q3v
532	Q1r	CF3	H	Q3w
533	Q1r	CF3	H	Q3x
534	Q1r	CF3	H	Q3y
535	Q1r	CF3	H	Q3z
536	Q1r	CF3	Me	Q3v
537	Q1r	CF3	Me	Q3w

538	Q1r	CF3	Me	Q3x
539	Q1r	CF3	Me	Q3y
540	Q1r	CF3	Me	Q3z
541	Q1s	H	H	Q3v
542	Q1s	H	H	Q3w
543	Q1s	H	H	Q3x
544	Q1s	H	H	Q3y
545	Q1s	H	H	Q3z
546	Q1s	H	Me	Q3v
547	Q1s	H	Me	Q3w
548	Q1s	H	Me	Q3x
549	Q1s	H	Me	Q3y
550	Q1s	H	Me	Q3z
551	Q1s	Me	H	Q3v
552	Q1s	Me	H	Q3w
553	Q1s	Me	H	Q3x
554	Q1s	Me	H	Q3y
555	Q1s	Me	H	Q3z
556	Q1s	Me	Me	Q3v
557	Q1s	Me	Me	Q3w
558	Q1s	Me	Me	Q3x
559	Q1s	Me	Me	Q3y
560	Q1s	Me	Me	Q3z
561	Q1s	CF3	H	Q3v
562	Q1s	CF3	H	Q3w
563	Q1s	CF3	H	Q3x
564	Q1s	CF3	H	Q3y
565	Q1s	CF3	H	Q3z
566	Q1s	CF3	Me	Q3v
567	Q1s	CF3	Me	Q3w
568	Q1s	CF3	Me	Q3x
569	Q1s	CF3	Me	Q3y
570	Q1s	CF3	Me	Q3z
571	Q1t	H	H	Q3v
572	Q1t	H	H	Q3w
573	Q1t	H	H	Q3x

574	Q1t	H	H	Q3y
575	Q1t	H	H	Q3z
576	Q1t	H	Me	Q3v
577	Q1t	H	Me	Q3w
578	Q1t	H	Me	Q3x
579	Q1t	H	Me	Q3y
580	Q1t	H	Me	Q3z
581	Q1t	Me	H	Q3v
582	Q1t	Me	H	Q3w
583	Q1t	Me	H	Q3x
584	Q1t	Me	H	Q3y
585	Q1t	Me	H	Q3z
586	Q1t	Me	Me	Q3v
587	Q1t	Me	Me	Q3w
588	Q1t	Me	Me	Q3x
589	Q1t	Me	Me	Q3y
590	Q1t	Me	Me	Q3z
591	Q1t	CF3	H	Q3v
592	Q1t	CF3	H	Q3w
593	Q1t	CF3	H	Q3x
594	Q1t	CF3	H	Q3y
595	Q1t	CF3	H	Q3z
596	Q1t	CF3	Me	Q3v
597	Q1t	CF3	Me	Q3w
598	Q1t	CF3	Me	Q3x
599	Q1t	CF3	Me	Q3y
600	Q1t	CF3	Me	Q3z
601	Q1u	H	H	Q3v
602	Q1u	H	H	Q3w
603	Q1u	H	H	Q3x
604	Q1u	H	H	Q3y
605	Q1u	H	H	Q3z
606	Q1u	H	Me	Q3v
607	Q1u	H	Me	Q3w
608	Q1u	H	Me	Q3x
609	Q1u	H	Me	Q3y

610	Q1u	H	Me	Q3z
611	Q1u	Me	H	Q3v
612	Q1u	Me	H	Q3w
613	Q1u	Me	H	Q3x
614	Q1u	Me	H	Q3y
615	Q1u	Me	H	Q3z
616	Q1u	Me	Me	Q3v
617	Q1u	Me	Me	Q3w
618	Q1u	Me	Me	Q3x
619	Q1u	Me	Me	Q3y
620	Q1u	Me	Me	Q3z
621	Q1u	CF3	H	Q3v
622	Q1u	CF3	H	Q3w
623	Q1u	CF3	H	Q3x
624	Q1u	CF3	H	Q3y
625	Q1u	CF3	H	Q3z
626	Q1u	CF3	Me	Q3v
627	Q1u	CF3	Me	Q3w
628	Q1u	CF3	Me	Q3x
629	Q1u	CF3	Me	Q3y
630	Q1u	CF3	Me	Q3z
631	Q1v	H	H	Q3v
632	Q1v	H	H	Q3w
633	Q1v	H	H	Q3x
634	Q1v	H	H	Q3y
635	Q1v	H	H	Q3z
636	Q1v	H	Me	Q3v
637	Q1v	H	Me	Q3w
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639	Q1v	H	Me	Q3y
640	Q1v	H	Me	Q3z
641	Q1v	Me	H	Q3v
642	Q1v	Me	H	Q3w
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644	Q1v	Me	H	Q3y
645	Q1v	Me	H	Q3z

646	Q1v	Me	Me	Q3v
647	Q1v	Me	Me	Q3w
648	Q1v	Me	Me	Q3x
649	Q1v	Me	Me	Q3y
650	Q1v	Me	Me	Q3z
651	Q1v	CF3	H	Q3v
652	Q1v	CF3	H	Q3w
653	Q1v	CF3	H	Q3x
654	Q1v	CF3	H	Q3y
655	Q1v	CF3	H	Q3z
656	Q1v	CF3	Me	Q3v
657	Q1v	CF3	Me	Q3w
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659	Q1v	CF3	Me	Q3y
660	Q1v	CF3	Me	Q3z
661	Q1w	H	H	Q3v
662	Q1w	H	H	Q3w
663	Q1w	H	H	Q3x
664	Q1w	H	H	Q3y
665	Q1w	H	H	Q3z
666	Q1w	H	Me	Q3v
667	Q1w	H	Me	Q3w
668	Q1w	H	Me	Q3x
669	Q1w	H	Me	Q3y
670	Q1w	H	Me	Q3z
671	Q1w	Me	H	Q3v
672	Q1w	Me	H	Q3w
673	Q1w	Me	H	Q3x
674	Q1w	Me	H	Q3y
675	Q1w	Me	H	Q3z
676	Q1w	Me	Me	Q3v
677	Q1w	Me	Me	Q3w
678	Q1w	Me	Me	Q3x
679	Q1w	Me	Me	Q3y
680	Q1w	Me	Me	Q3z
681	Q1w	CF3	H	Q3v

682	Q1w	CF3	H	Q3w
683	Q1w	CF3	H	Q3x
684	Q1w	CF3	H	Q3y
685	Q1w	CF3	H	Q3z
686	Q1w	CF3	Me	Q3v
687	Q1w	CF3	Me	Q3w
688	Q1w	CF3	Me	Q3x
689	Q1w	CF3	Me	Q3y
690	Q1w	CF3	Me	Q3z
691	Q1x	H	H	Q3v
692	Q1x	H	H	Q3w
693	Q1x	H	H	Q3x
694	Q1x	H	H	Q3y
695	Q1x	H	H	Q3z
696	Q1x	H	Me	Q3v
697	Q1x	H	Me	Q3w
698	Q1x	H	Me	Q3x
699	Q1x	H	Me	Q3y
700	Q1x	H	Me	Q3z
701	Q1x	Me	H	Q3v
702	Q1x	Me	H	Q3w
703	Q1x	Me	H	Q3x
704	Q1x	Me	H	Q3y
705	Q1x	Me	H	Q3z
706	Q1x	Me	Me	Q3v
707	Q1x	Me	Me	Q3w
708	Q1x	Me	Me	Q3x
709	Q1x	Me	Me	Q3y
710	Q1x	Me	Me	Q3z
711	Q1x	CF3	H	Q3v
712	Q1x	CF3	H	Q3w
713	Q1x	CF3	H	Q3x
714	Q1x	CF3	H	Q3y
715	Q1x	CF3	H	Q3z
716	Q1x	CF3	Me	Q3v
717	Q1x	CF3	Me	Q3w

718	Q1x	CF3	Me	Q3x
719	Q1x	CF3	Me	Q3y
720	Q1x	CF3	Me	Q3z
721	Q1y	H	H	Q3v
722	Q1y	H	H	Q3w
723	Q1y	H	H	Q3x
724	Q1y	H	H	Q3y
725	Q1y	H	H	Q3z
726	Q1y	H	Me	Q3v
727	Q1y	H	Me	Q3w
728	Q1y	H	Me	Q3x
729	Q1y	H	Me	Q3y
730	Q1y	H	Me	Q3z
731	Q1y	Me	H	Q3v
732	Q1y	Me	H	Q3w
733	Q1y	Me	H	Q3x
734	Q1y	Me	H	Q3y
735	Q1y	Me	H	Q3z
736	Q1y	Me	Me	Q3v
737	Q1y	Me	Me	Q3w
738	Q1y	Me	Me	Q3x
739	Q1y	Me	Me	Q3y
740	Q1y	Me	Me	Q3z
741	Q1y	CF3	H	Q3v
742	Q1y	CF3	H	Q3w
743	Q1y	CF3	H	Q3x
744	Q1y	CF3	H	Q3y
745	Q1y	CF3	H	Q3z
746	Q1y	CF3	Me	Q3v
747	Q1y	CF3	Me	Q3w
748	Q1y	CF3	Me	Q3x
749	Q1y	CF3	Me	Q3y
750	Q1y	CF3	Me	Q3z
751	Q1z	H	H	Q3v
752	Q1z	H	H	Q3w
753	Q1z	H	H	Q3x

754	Q1z	H	H	Q3y
755	Q1z	H	H	Q3z
756	Q1z	H	Me	Q3v
757	Q1z	H	Me	Q3w
758	Q1z	H	Me	Q3x
759	Q1z	H	Me	Q3y
760	Q1z	H	Me	Q3z
761	Q1z	Me	H	Q3v
762	Q1z	Me	H	Q3w
763	Q1z	Me	H	Q3x
764	Q1z	Me	H	Q3y
765	Q1z	Me	H	Q3z
766	Q1z	Me	Me	Q3v
767	Q1z	Me	Me	Q3w
768	Q1z	Me	Me	Q3x
769	Q1z	Me	Me	Q3y
770	Q1z	Me	Me	Q3z
771	Q1z	CF3	H	Q3v
772	Q1z	CF3	H	Q3w
773	Q1z	CF3	H	Q3x
774	Q1z	CF3	H	Q3y
775	Q1z	CF3	H	Q3z
776	Q1z	CF3	Me	Q3v
777	Q1z	CF3	Me	Q3w
778	Q1z	CF3	Me	Q3x
779	Q1z	CF3	Me	Q3y
780	Q1z	CF3	Me	Q3z

132) The compounds wherein R⁷, R⁸, R⁹ and R¹⁰ are any of the following combinations in Table 3, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof. The symbols in Table 3
5 denote the following substituents.

【Ka 12】

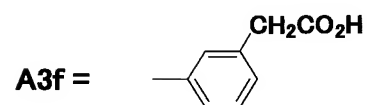
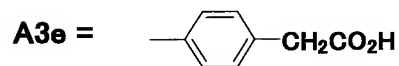
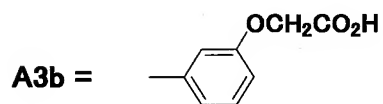
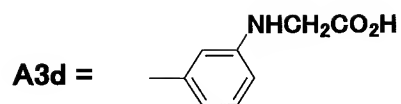
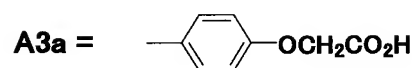
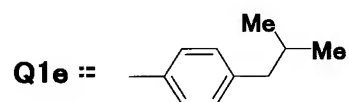
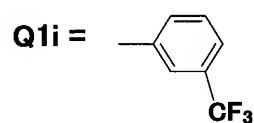
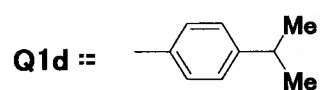
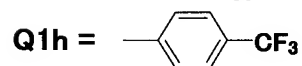
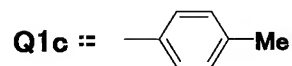
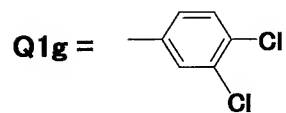
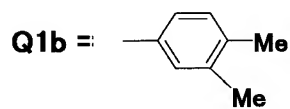
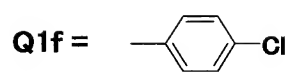
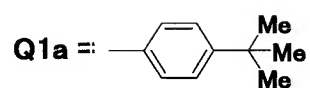


Table 3

No	R ⁷	R ⁸	R ⁹	R ¹⁰
1	Q1a	Me	Me	A3a
2	Q1a	Me	Me	A3b
3	Q1a	Me	Me	A3c
4	Q1a	Me	Me	A3d
5	Q1a	Me	Me	A3e
6	Q1a	Me	Me	A3f
7	Q1a	Me	H	A3a
8	Q1a	Me	H	A3b
9	Q1a	Me	H	A3c
10	Q1a	Me	H	A3d
11	Q1a	Me	H	A3e
12	Q1a	Me	H	A3f
13	Q1a	CF3	Me	A3a
14	Q1a	CF3	Me	A3b
15	Q1a	CF3	Me	A3c
16	Q1a	CF3	Me	A3d
17	Q1a	CF3	Me	A3e
18	Q1a	CF3	Me	A3f
19	Q1a	CF3	H	A3a
20	Q1a	CF3	H	A3b
21	Q1a	CF3	H	A3c
22	Q1a	CF3	H	A3d
23	Q1a	CF3	H	A3e
24	Q1a	CF3	H	A3f
25	Q1b	Me	Me	A3a
26	Q1b	Me	Me	A3b
27	Q1b	Me	Me	A3c
28	Q1b	Me	Me	A3d
29	Q1b	Me	Me	A3e
30	Q1b	Me	Me	A3f
31	Q1b	Me	H	A3a
32	Q1b	Me	H	A3b
33	Q1b	Me	H	A3c

34	Q1b	Me	H	A3d
35	Q1b	Me	H	A3e
36	Q1b	Me	H	A3f
37	Q1b	CF3	Me	A3a
38	Q1b	CF3	Me	A3b
39	Q1b	CF3	Me	A3c
40	Q1b	CF3	Me	A3d
41	Q1b	CF3	Me	A3e
42	Q1b	CF3	Me	A3f
43	Q1b	CF3	H	A3a
44	Q1b	CF3	H	A3b
45	Q1b	CF3	H	A3c
46	Q1b	CF3	H	A3d
47	Q1b	CF3	H	A3e
48	Q1b	CF3	H	A3f
49	Q1c	Me	Me	A3a
50	Q1c	Me	Me	A3b
51	Q1c	Me	Me	A3c
52	Q1c	Me	Me	A3d
53	Q1c	Me	Me	A3e
54	Q1c	Me	Me	A3f
55	Q1c	Me	H	A3a
56	Q1c	Me	H	A3b
57	Q1c	Me	H	A3c
58	Q1c	Me	H	A3d
59	Q1c	Me	H	A3e
60	Q1c	Me	H	A3f
61	Q1c	CF3	Me	A3a
62	Q1c	CF3	Me	A3b
63	Q1c	CF3	Me	A3c
64	Q1c	CF3	Me	A3d
65	Q1c	CF3	Me	A3e
66	Q1c	CF3	Me	A3f
67	Q1c	CF3	H	A3a
68	Q1c	CF3	H	A3b
69	Q1c	CF3	H	A3c

70	Q1c	CF3	H	A3d
71	Q1c	CF3	H	A3e
72	Q1c	CF3	H	A3f
73	Q1d	Me	Me	A3a
74	Q1d	Me	Me	A3b
75	Q1d	Me	Me	A3c
76	Q1d	Me	Me	A3d
77	Q1d	Me	Me	A3e
78	Q1d	Me	Me	A3f
79	Q1d	Me	H	A3a
80	Q1d	Me	H	A3b
81	Q1d	Me	H	A3c
82	Q1d	Me	H	A3d
83	Q1d	Me	H	A3e
84	Q1d	Me	H	A3f
85	Q1d	CF3	Me	A3a
86	Q1d	CF3	Me	A3b
87	Q1d	CF3	Me	A3c
88	Q1d	CF3	Me	A3d
89	Q1d	CF3	Me	A3e
90	Q1d	CF3	Me	A3f
91	Q1d	CF3	H	A3a
92	Q1d	CF3	H	A3b
93	Q1d	CF3	H	A3c
94	Q1d	CF3	H	A3d
95	Q1d	CF3	H	A3e
96	Q1d	CF3	H	A3f
97	Q1e	Me	Me	A3a
98	Q1e	Me	Me	A3b
99	Q1e	Me	Me	A3c
100	Q1e	Me	Me	A3d
101	Q1e	Me	Me	A3e
102	Q1e	Me	Me	A3f
103	Q1e	Me	H	A3a
104	Q1e	Me	H	A3b
105	Q1e	Me	H	A3c

106	Q1e	Me	H	A3d
107	Q1e	Me	H	A3e
108	Q1e	Me	H	A3f
109	Q1e	CF3	Me	A3a
110	Q1e	CF3	Me	A3b
111	Q1e	CF3	Me	A3c
112	Q1e	CF3	Me	A3d
113	Q1e	CF3	Me	A3e
114	Q1e	CF3	Me	A3f
115	Q1e	CF3	H	A3a
116	Q1e	CF3	H	A3b
117	Q1e	CF3	H	A3c
118	Q1e	CF3	H	A3d
119	Q1e	CF3	H	A3e
120	Q1e	CF3	H	A3f
121	Q1f	Me	Me	A3a
122	Q1f	Me	Me	A3b
123	Q1f	Me	Me	A3c
124	Q1f	Me	Me	A3d
125	Q1f	Me	Me	A3e
126	Q1f	Me	Me	A3f
127	Q1f	Me	H	A3a
128	Q1f	Me	H	A3b
129	Q1f	Me	H	A3c
130	Q1f	Me	H	A3d
131	Q1f	Me	H	A3e
132	Q1f	Me	H	A3f
133	Q1f	CF3	Me	A3a
134	Q1f	CF3	Me	A3b
135	Q1f	CF3	Me	A3c
136	Q1f	CF3	Me	A3d
137	Q1f	CF3	Me	A3e
138	Q1f	CF3	Me	A3f
139	Q1f	CF3	H	A3a
140	Q1f	CF3	H	A3b
141	Q1f	CF3	H	A3c

142	Q1f	CF3	H	A3d
143	Q1f	CF3	H	A3e
144	Q1f	CF3	H	A3f
145	Q1g	Me	Me	A3a
146	Q1g	Me	Me	A3b
147	Q1g	Me	Me	A3c
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149	Q1g	Me	Me	A3e
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151	Q1g	Me	H	A3a
152	Q1g	Me	H	A3b
153	Q1g	Me	H	A3c
154	Q1g	Me	H	A3d
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165	Q1g	CF3	H	A3c
166	Q1g	CF3	H	A3d
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175	Q1h	Me	H	A3a
176	Q1h	Me	H	A3b
177	Q1h	Me	H	A3c

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182	Q1h	CF3	Me	A3b
183	Q1h	CF3	Me	A3c
184	Q1h	CF3	Me	A3d
185	Q1h	CF3	Me	A3e
186	Q1h	CF3	Me	A3f
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189	Q1h	CF3	H	A3c
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194	Q1i	Me	Me	A3b
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196	Q1i	Me	Me	A3d
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199	Q1i	Me	H	A3a
200	Q1i	Me	H	A3b
201	Q1i	Me	H	A3c
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211	Q1i	CF3	H	A3a
212	Q1i	CF3	H	A3b
213	Q1i	CF3	H	A3c

214	Q1i	CF3	H	A3d
215	Q1i	CF3	H	A3e
216	Q1i	CF3	H	A3f

133) The compounds wherein R^{12} , R^{13} , R^{14} and R^{15} are any of the following combinations in Table 4, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof. The symbols in Table 4
5 denote the following substituents.

【Ka 13】

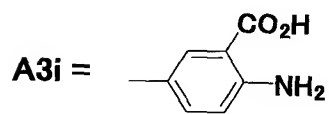
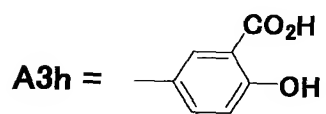
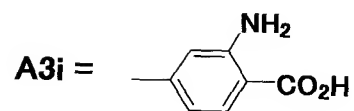
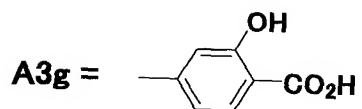
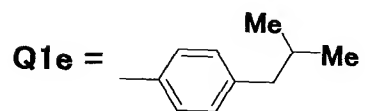
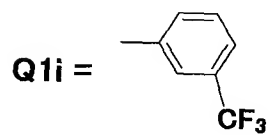
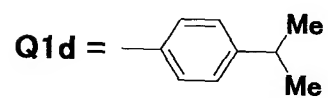
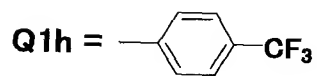
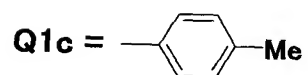
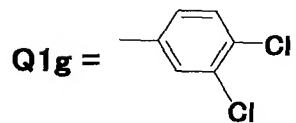
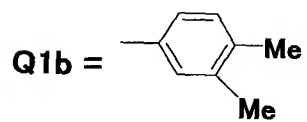
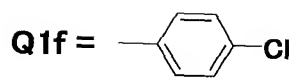
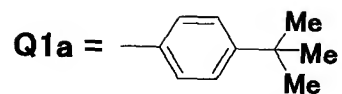


Table 4

No	R ¹²	R ¹³	R ¹⁴	R ¹⁵
1	Q1a	Me	H	A3g
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3	Q1a	Me	H	A3i
4	Q1a	Me	H	A3j
5	Q1a	Me	Me	A3g
6	Q1a	Me	Me	A3h
7	Q1a	Me	Me	A3i
8	Q1a	Me	Me	A3j
9	Q1a	CF3	H	A3g
10	Q1a	CF3	H	A3h
11	Q1a	CF3	H	A3i
12	Q1a	CF3	H	A3j
13	Q1a	CF3	Me	A3g
14	Q1a	CF3	Me	A3h
15	Q1a	CF3	Me	A3i
16	Q1a	CF3	Me	A3j
17	Q1b	Me	H	A3g
18	Q1b	Me	H	A3h
19	Q1b	Me	H	A3i
20	Q1b	Me	H	A3j
21	Q1b	Me	Me	A3g
22	Q1b	Me	Me	A3h
23	Q1b	Me	Me	A3i
24	Q1b	Me	Me	A3j
25	Q1b	CF3	H	A3g
26	Q1b	CF3	H	A3h
27	Q1b	CF3	H	A3i
28	Q1b	CF3	H	A3j
29	Q1b	CF3	Me	A3g
30	Q1b	CF3	Me	A3h
31	Q1b	CF3	Me	A3i
32	Q1b	CF3	Me	A3j
33	Q1c	Me	H	A3g

34	Q1c	Me	H	A3h
35	Q1c	Me	H	A3i
36	Q1c	Me	H	A3j
37	Q1c	Me	Me	A3g
38	Q1c	Me	Me	A3h
39	Q1c	Me	Me	A3i
40	Q1c	Me	Me	A3j
41	Q1c	CF3	H	A3g
42	Q1c	CF3	H	A3h
43	Q1c	CF3	H	A3i
44	Q1c	CF3	H	A3j
45	Q1c	CF3	Me	A3g
46	Q1c	CF3	Me	A3h
47	Q1c	CF3	Me	A3i
48	Q1c	CF3	Me	A3j
49	Q1d	Me	H	A3g
50	Q1d	Me	H	A3h
51	Q1d	Me	H	A3i
52	Q1d	Me	H	A3j
53	Q1d	Me	Me	A3g
54	Q1d	Me	Me	A3h
55	Q1d	Me	Me	A3i
56	Q1d	Me	Me	A3j
57	Q1d	CF3	H	A3g
58	Q1d	CF3	H	A3h
59	Q1d	CF3	H	A3i
60	Q1d	CF3	H	A3j
61	Q1d	CF3	Me	A3g
62	Q1d	CF3	Me	A3h
63	Q1d	CF3	Me	A3i
64	Q1d	CF3	Me	A3j
65	Q1e	Me	H	A3g
66	Q1e	Me	H	A3h
67	Q1e	Me	H	A3i
68	Q1e	Me	H	A3j
69	Q1e	Me	Me	A3g

70	Q1e	Me	Me	A3h
71	Q1e	Me	Me	A3i
72	Q1e	Me	Me	A3j
73	Q1e	CF3	H	A3g
74	Q1e	CF3	H	A3h
75	Q1e	CF3	H	A3i
76	Q1e	CF3	H	A3j
77	Q1e	CF3	Me	A3g
78	Q1e	CF3	Me	A3h
79	Q1e	CF3	Me	A3i
80	Q1e	CF3	Me	A3j
81	Q1f	Me	H	A3g
82	Q1f	Me	H	A3h
83	Q1f	Me	H	A3i
84	Q1f	Me	H	A3j
85	Q1f	Me	Me	A3g
86	Q1f	Me	Me	A3h
87	Q1f	Me	Me	A3i
88	Q1f	Me	Me	A3j
89	Q1f	CF3	H	A3g
90	Q1f	CF3	H	A3h
91	Q1f	CF3	H	A3i
92	Q1f	CF3	H	A3j
93	Q1f	CF3	Me	A3g
94	Q1f	CF3	Me	A3h
95	Q1f	CF3	Me	A3i
96	Q1f	CF3	Me	A3j
97	Q1g	Me	H	A3g
98	Q1g	Me	H	A3h
99	Q1g	Me	H	A3i
100	Q1g	Me	H	A3j
101	Q1g	Me	Me	A3g
102	Q1g	Me	Me	A3h
103	Q1g	Me	Me	A3i
104	Q1g	Me	Me	A3j
105	Q1g	CF3	H	A3g

106	Q1g	CF3	H	A3h
107	Q1g	CF3	H	A3i
108	Q1g	CF3	H	A3j
109	Q1g	CF3	Me	A3g
110	Q1g	CF3	Me	A3h
111	Q1g	CF3	Me	A3i
112	Q1g	CF3	Me	A3j
113	Q1h	Me	H	A3g
114	Q1h	Me	H	A3h
115	Q1h	Me	H	A3i
116	Q1h	Me	H	A3j
117	Q1h	Me	Me	A3g
118	Q1h	Me	Me	A3h
119	Q1h	Me	Me	A3i
120	Q1h	Me	Me	A3j
121	Q1h	CF3	H	A3g
122	Q1h	CF3	H	A3h
123	Q1h	CF3	H	A3i
124	Q1h	CF3	H	A3j
125	Q1h	CF3	Me	A3g
126	Q1h	CF3	Me	A3h
127	Q1h	CF3	Me	A3i
128	Q1h	CF3	Me	A3j
129	Q1i	Me	H	A3g
130	Q1i	Me	H	A3h
131	Q1i	Me	H	A3i
132	Q1i	Me	H	A3j
133	Q1i	Me	Me	A3g
134	Q1i	Me	Me	A3h
135	Q1i	Me	Me	A3i
136	Q1i	Me	Me	A3j
137	Q1i	CF3	H	A3g
138	Q1i	CF3	H	A3h
139	Q1i	CF3	H	A3i
140	Q1i	CF3	H	A3j
141	Q1i	CF3	Me	A3g

142	Q1i	CF3	Me	A3h
143	Q1i	CF3	Me	A3i
144	Q1i	CF3	Me	A3j

134) The compounds wherein R⁷, R⁸, R⁹ and R¹⁰ are any of the following combinations in Table 5, tautomers, prodrugs or pharmaceutically acceptable salts of the compounds or solvates thereof. The symbols in Table 5

5 denote the following substituents.

【Ka 14】

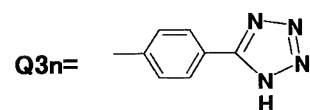
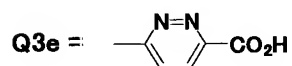
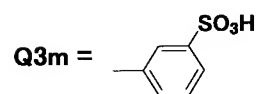
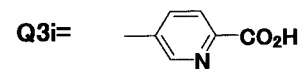
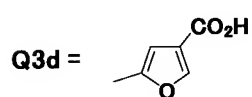
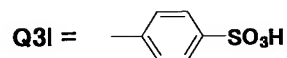
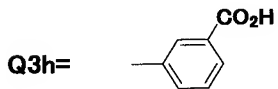
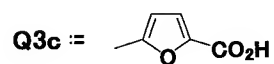
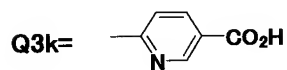
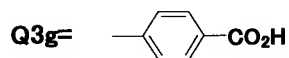
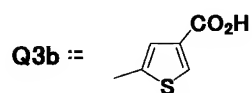
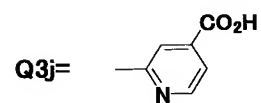
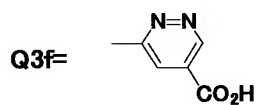
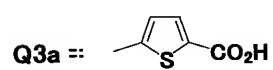
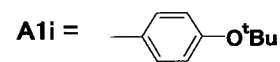
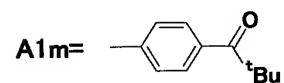
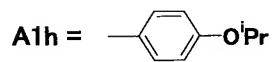
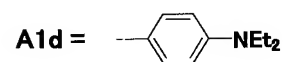
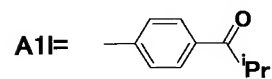
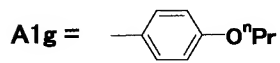
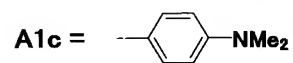
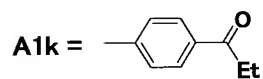
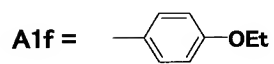
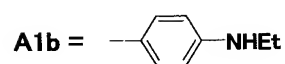
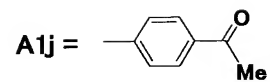
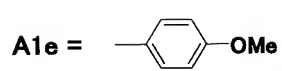
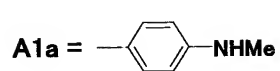


Table 5

	R ⁷	R ⁸	R ⁹	R ¹⁰
1	A1a	Me	H	Q3a
2	A1a	Me	H	Q3b
3	A1a	Me	H	Q3c
4	A1a	Me	H	Q3d
5	A1a	Me	H	Q3e
6	A1a	Me	H	Q3f
7	A1a	Me	H	Q3g
8	A1a	Me	H	Q3h
9	A1a	Me	H	Q3i
10	A1a	Me	H	Q3j
11	A1a	Me	H	Q3k
12	A1a	Me	H	Q3l
13	A1a	Me	H	Q3m
14	A1a	Me	H	Q3n
15	A1a	Me	Me	Q3a
16	A1a	Me	Me	Q3b
17	A1a	Me	Me	Q3c
18	A1a	Me	Me	Q3d
19	A1a	Me	Me	Q3e
20	A1a	Me	Me	Q3f
21	A1a	Me	Me	Q3g
22	A1a	Me	Me	Q3h
23	A1a	Me	Me	Q3i
24	A1a	Me	Me	Q3j
25	A1a	Me	Me	Q3k
26	A1a	Me	Me	Q3l
27	A1a	Me	Me	Q3m
28	A1a	Me	Me	Q3n
29	A1b	Me	H	Q3a
30	A1b	Me	H	Q3b
31	A1b	Me	H	Q3c
32	A1b	Me	H	Q3d
33	A1b	Me	H	Q3e

34	A1b	Me	H	Q3f
35	A1b	Me	H	Q3g
36	A1b	Me	H	Q3h
37	A1b	Me	H	Q3i
38	A1b	Me	H	Q3j
39	A1b	Me	H	Q3k
40	A1b	Me	H	Q3l
41	A1b	Me	H	Q3m
42	A1b	Me	H	Q3n
43	A1b	Me	Me	Q3a
44	A1b	Me	Me	Q3b
45	A1b	Me	Me	Q3c
46	A1b	Me	Me	Q3d
47	A1b	Me	Me	Q3e
48	A1b	Me	Me	Q3f
49	A1b	Me	Me	Q3g
50	A1b	Me	Me	Q3h
51	A1b	Me	Me	Q3i
52	A1b	Me	Me	Q3j
53	A1b	Me	Me	Q3k
54	A1b	Me	Me	Q3l
55	A1b	Me	Me	Q3m
56	A1b	Me	Me	Q3n
57	A1c	Me	H	Q3a
58	A1c	Me	H	Q3b
59	A1c	Me	H	Q3c
60	A1c	Me	H	Q3d
61	A1c	Me	H	Q3e
62	A1c	Me	H	Q3f
63	A1c	Me	H	Q3g
64	A1c	Me	H	Q3h
65	A1c	Me	H	Q3i
66	A1c	Me	H	Q3j
67	A1c	Me	H	Q3k
68	A1c	Me	H	Q3l
69	A1c	Me	H	Q3m

70	A1c	Me	H	Q3n
71	A1c	Me	Me	Q3a
72	A1c	Me	Me	Q3b
73	A1c	Me	Me	Q3c
74	A1c	Me	Me	Q3d
75	A1c	Me	Me	Q3e
76	A1c	Me	Me	Q3f
77	A1c	Me	Me	Q3g
78	A1c	Me	Me	Q3h
79	A1c	Me	Me	Q3i
80	A1c	Me	Me	Q3j
81	A1c	Me	Me	Q3k
82	A1c	Me	Me	Q3l
83	A1c	Me	Me	Q3m
84	A1c	Me	Me	Q3n
85	A1d	Me	H	Q3a
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89	A1d	Me	H	Q3e
90	A1d	Me	H	Q3f
91	A1d	Me	H	Q3g
92	A1d	Me	H	Q3h
93	A1d	Me	H	Q3i
94	A1d	Me	H	Q3j
95	A1d	Me	H	Q3k
96	A1d	Me	H	Q3l
97	A1d	Me	H	Q3m
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110	A1d	Me	Me	Q3l
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112	A1d	Me	Me	Q3n
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126	A1e	Me	H	Q3n
127	A1e	Me	Me	Q3a
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132	A1e	Me	Me	Q3f
133	A1e	Me	Me	Q3g
134	A1e	Me	Me	Q3h
135	A1e	Me	Me	Q3i
136	A1e	Me	Me	Q3j
137	A1e	Me	Me	Q3k
138	A1e	Me	Me	Q3l
139	A1e	Me	Me	Q3m
140	A1e	Me	Me	Q3n
141	A1f	Me	H	Q3a

142	A1f	Me	H	Q3b
143	A1f	Me	H	Q3c
144	A1f	Me	H	Q3d
145	A1f	Me	H	Q3e
146	A1f	Me	H	Q3f
147	A1f	Me	H	Q3g
148	A1f	Me	H	Q3h
149	A1f	Me	H	Q3i
150	A1f	Me	H	Q3j
151	A1f	Me	H	Q3k
152	A1f	Me	H	Q3l
153	A1f	Me	H	Q3m
154	A1f	Me	H	Q3n
155	A1f	Me	Me	Q3a
156	A1f	Me	Me	Q3b
157	A1f	Me	Me	Q3c
158	A1f	Me	Me	Q3d
159	A1f	Me	Me	Q3e
160	A1f	Me	Me	Q3f
161	A1f	Me	Me	Q3g
162	A1f	Me	Me	Q3h
163	A1f	Me	Me	Q3i
164	A1f	Me	Me	Q3j
165	A1f	Me	Me	Q3k
166	A1f	Me	Me	Q3l
167	A1f	Me	Me	Q3m
168	A1f	Me	Me	Q3n
169	A1g	Me	H	Q3a
170	A1g	Me	H	Q3b
171	A1g	Me	H	Q3c
172	A1g	Me	H	Q3d
173	A1g	Me	H	Q3e
174	A1g	Me	H	Q3f
175	A1g	Me	H	Q3g
176	A1g	Me	H	Q3h
177	A1g	Me	H	Q3i

178	A1g	Me	H	Q3j
179	A1g	Me	H	Q3k
180	A1g	Me	H	Q3l
181	A1g	Me	H	Q3m
182	A1g	Me	H	Q3n
183	A1g	Me	Me	Q3a
184	A1g	Me	Me	Q3b
185	A1g	Me	Me	Q3c
186	A1g	Me	Me	Q3d
187	A1g	Me	Me	Q3e
188	A1g	Me	Me	Q3f
189	A1g	Me	Me	Q3g
190	A1g	Me	Me	Q3h
191	A1g	Me	Me	Q3i
192	A1g	Me	Me	Q3j
193	A1g	Me	Me	Q3k
194	A1g	Me	Me	Q3l
195	A1g	Me	Me	Q3m
196	A1g	Me	Me	Q3n
197	A1h	Me	H	Q3a
198	A1h	Me	H	Q3b
199	A1h	Me	H	Q3c
200	A1h	Me	H	Q3d
201	A1h	Me	H	Q3e
202	A1h	Me	H	Q3f
203	A1h	Me	H	Q3g
204	A1h	Me	H	Q3h
205	A1h	Me	H	Q3i
206	A1h	Me	H	Q3j
207	A1h	Me	H	Q3k
208	A1h	Me	H	Q3l
209	A1h	Me	H	Q3m
210	A1h	Me	H	Q3n
211	A1h	Me	Me	Q3a
212	A1h	Me	Me	Q3b
213	A1h	Me	Me	Q3c

214	A1h	Me	Me	Q3d
215	A1h	Me	Me	Q3e
216	A1h	Me	Me	Q3f
217	A1h	Me	Me	Q3g
218	A1h	Me	Me	Q3h
219	A1h	Me	Me	Q3i
220	A1h	Me	Me	Q3j
221	A1h	Me	Me	Q3k
222	A1h	Me	Me	Q3l
223	A1h	Me	Me	Q3m
224	A1h	Me	Me	Q3n
225	A1i	Me	H	Q3a
226	A1i	Me	H	Q3b
227	A1i	Me	H	Q3c
228	A1i	Me	H	Q3d
229	A1i	Me	H	Q3e
230	A1i	Me	H	Q3f
231	A1i	Me	H	Q3g
232	A1i	Me	H	Q3h
233	A1i	Me	H	Q3i
234	A1i	Me	H	Q3j
235	A1i	Me	H	Q3k
236	A1i	Me	H	Q3l
237	A1i	Me	H	Q3m
238	A1i	Me	H	Q3n
239	A1i	Me	Me	Q3a
240	A1i	Me	Me	Q3b
241	A1i	Me	Me	Q3c
242	A1i	Me	Me	Q3d
243	A1i	Me	Me	Q3e
244	A1i	Me	Me	Q3f
245	A1i	Me	Me	Q3g
246	A1i	Me	Me	Q3h
247	A1i	Me	Me	Q3i
248	A1i	Me	Me	Q3j
249	A1i	Me	Me	Q3k

250	A1i	Me	Me	Q3l
251	A1i	Me	Me	Q3m
252	A1i	Me	Me	Q3n
253	A1j	Me	H	Q3a
254	A1j	Me	H	Q3b
255	A1j	Me	H	Q3c
256	A1j	Me	H	Q3d
257	A1j	Me	H	Q3e
258	A1j	Me	H	Q3f
259	A1j	Me	H	Q3g
260	A1j	Me	H	Q3h
261	A1j	Me	H	Q3i
262	A1j	Me	H	Q3j
263	A1j	Me	H	Q3k
264	A1j	Me	H	Q3l
265	A1j	Me	H	Q3m
266	A1j	Me	H	Q3n
267	A1j	Me	Me	Q3a
268	A1j	Me	Me	Q3b
269	A1j	Me	Me	Q3c
270	A1j	Me	Me	Q3d
271	A1j	Me	Me	Q3e
272	A1j	Me	Me	Q3f
273	A1j	Me	Me	Q3g
274	A1j	Me	Me	Q3h
275	A1j	Me	Me	Q3i
276	A1j	Me	Me	Q3j
277	A1j	Me	Me	Q3k
278	A1j	Me	Me	Q3l
279	A1j	Me	Me	Q3m
280	A1j	Me	Me	Q3n
281	A1k	Me	H	Q3a
282	A1k	Me	H	Q3b
283	A1k	Me	H	Q3c
284	A1k	Me	H	Q3d
285	A1k	Me	H	Q3e

286	A1k	Me	H	Q3f
287	A1k	Me	H	Q3g
288	A1k	Me	H	Q3h
289	A1k	Me	H	Q3i
290	A1k	Me	H	Q3j
291	A1k	Me	H	Q3k
292	A1k	Me	H	Q3l
293	A1k	Me	H	Q3m
294	A1k	Me	H	Q3n
295	A1k	Me	Me	Q3a
296	A1k	Me	Me	Q3b
297	A1k	Me	Me	Q3c
298	A1k	Me	Me	Q3d
299	A1k	Me	Me	Q3e
300	A1k	Me	Me	Q3f
301	A1k	Me	Me	Q3g
302	A1k	Me	Me	Q3h
303	A1k	Me	Me	Q3i
304	A1k	Me	Me	Q3j
305	A1k	Me	Me	Q3k
306	A1k	Me	Me	Q3l
307	A1k	Me	Me	Q3m
308	A1k	Me	Me	Q3n
309	A1l	Me	H	Q3a
310	A1l	Me	H	Q3b
311	A1l	Me	H	Q3c
312	A1l	Me	H	Q3d
313	A1l	Me	H	Q3e
314	A1l	Me	H	Q3f
315	A1l	Me	H	Q3g
316	A1l	Me	H	Q3h
317	A1l	Me	H	Q3i
318	A1l	Me	H	Q3j
319	A1l	Me	H	Q3k
320	A1l	Me	H	Q3l
321	A1l	Me	H	Q3m

322	A1l	Me	H	Q3n
323	A1l	Me	Me	Q3a
324	A1l	Me	Me	Q3b
325	A1l	Me	Me	Q3c
326	A1l	Me	Me	Q3d
327	A1l	Me	Me	Q3e
328	A1l	Me	Me	Q3f
329	A1l	Me	Me	Q3g
330	A1l	Me	Me	Q3h
331	A1l	Me	Me	Q3i
332	A1l	Me	Me	Q3j
333	A1l	Me	Me	Q3k
334	A1l	Me	Me	Q3l
335	A1l	Me	Me	Q3m
336	A1l	Me	Me	Q3n
337	A1m	Me	H	Q3a
338	A1m	Me	H	Q3b
339	A1m	Me	H	Q3c
340	A1m	Me	H	Q3d
341	A1m	Me	H	Q3e
342	A1m	Me	H	Q3f
343	A1m	Me	H	Q3g
344	A1m	Me	H	Q3h
345	A1m	Me	H	Q3i
346	A1m	Me	H	Q3j
347	A1m	Me	H	Q3k
348	A1m	Me	H	Q3l
349	A1m	Me	H	Q3m
350	A1m	Me	H	Q3n
351	A1m	Me	Me	Q3a
352	A1m	Me	Me	Q3b
353	A1m	Me	Me	Q3c
354	A1m	Me	Me	Q3d
355	A1m	Me	Me	Q3e
356	A1m	Me	Me	Q3f
357	A1m	Me	Me	Q3g

358	A1m	Me	Me	Q3h
359	A1m	Me	Me	Q3i
360	A1m	Me	Me	Q3j
361	A1m	Me	Me	Q3k
362	A1m	Me	Me	Q3l
363	A1m	Me	Me	Q3m
364	A1m	Me	Me	Q3n

135) The thrombopoietin receptor activators represented by 1).

136) The thrombopoietin receptor activators represented by 2).

137) The thrombopoietin receptor activators represented by 3).

138) The thrombopoietin receptor activators represented by 4).

139) The thrombopoietin receptor activators represented by 5).

140) The thrombopoietin receptor activators represented by 6).

141) The thrombopoietin receptor activators represented by 7).

142) The thrombopoietin receptor activators represented by 8).

143) The thrombopoietin receptor activators represented by 9).

144) The thrombopoietin receptor activators represented by 10).

145) The thrombopoietin receptor activators represented by 11).

146) The thrombopoietin receptor activators represented by 12).

5 147) The thrombopoietin receptor activators represented by 13).

148) The thrombopoietin receptor activators represented by 14).

10 149) The thrombopoietin receptor activators represented by 15).

150) The thrombopoietin receptor activators represented by 16).

151) The thrombopoietin receptor activators represented by 17).

15 152) The thrombopoietin receptor activators represented by 18).

153) The thrombopoietin receptor activators represented by 19).

20 154) The thrombopoietin receptor activators represented by 20).

155) The thrombopoietin receptor activators represented by 21).

156) The thrombopoietin receptor activators represented by 22).

25 157) The thrombopoietin receptor activators represented by 23).

158) The thrombopoietin receptor activators represented

by 24) .

159) The thrombopoietin receptor activators represented
by 25) .

160) The thrombopoietin receptor activators represented
5 by 26) .

161) The thrombopoietin receptor activators represented
by 27) .

162) The thrombopoietin receptor activators represented
by 28) .

10 163) The thrombopoietin receptor activators represented
by 29) .

164) The thrombopoietin receptor activators represented
by 30) .

165) The thrombopoietin receptor activators represented
15 by 31) .

166) The thrombopoietin receptor activators represented
by 32) .

167) The thrombopoietin receptor activators represented
by 33) .

20 168) The thrombopoietin receptor activators represented
by 34) .

169) The thrombopoietin receptor activators represented
by 35) .

170) The thrombopoietin receptor activators represented
25 by 36) .

171) The thrombopoietin receptor activators represented
by 37) .

172) The thrombopoietin receptor activators represented
by 38).

173) The thrombopoietin receptor activators represented
by 39).

5 174) The thrombopoietin receptor activators represented
by 40).

175) The thrombopoietin receptor activators represented
by 41).

10 176) The thrombopoietin receptor activators represented
by 42).

177) The thrombopoietin receptor activators represented
by 43).

178) The thrombopoietin receptor activators represented
by 44).

15 179) The thrombopoietin receptor activators represented
by 45).

180) The thrombopoietin receptor activators represented
by 46).

20 181) The thrombopoietin receptor activators represented
by 47).

182) The thrombopoietin receptor activators represented
by 48).

183) The thrombopoietin receptor activators represented
by 49).

25 184) The thrombopoietin receptor activators represented
by 50).

185) The thrombopoietin receptor activators represented

by 51) .

186) The thrombopoietin receptor activators represented
by 52) .

187) The thrombopoietin receptor activators represented
5 by 53) .

188) The thrombopoietin receptor activators represented
by 54) .

189) The thrombopoietin receptor activators represented
by 55) .

10 190) The thrombopoietin receptor activators represented
by 56) .

191) The thrombopoietin receptor activators represented
by 57) .

192) The thrombopoietin receptor activators represented
15 by 58) .

193) The thrombopoietin receptor activators represented
by 59) .

194) The thrombopoietin receptor activators represented
by 60) .

20 195) The thrombopoietin receptor activators represented
by 61) .

196) The thrombopoietin receptor activators represented
by 62) .

197) The thrombopoietin receptor activators represented
25 by 63) .

198) The thrombopoietin receptor activators represented
by 64) .

199) The thrombopoietin receptor activators represented by 65).

200) The thrombopoietin receptor activators represented by 66).

5 201) The thrombopoietin receptor activators represented by 67).

202) The thrombopoietin receptor activators represented by 68).

10 203) The thrombopoietin receptor activators represented by 69).

204) The thrombopoietin receptor activators represented by 70).

205) The thrombopoietin receptor activators represented by 71).

15 206) The thrombopoietin receptor activators represented by 72).

207) The thrombopoietin receptor activators represented by 73).

20 208) The thrombopoietin receptor activators represented by 74).

209) The thrombopoietin receptor activators represented by 75).

210) The thrombopoietin receptor activators represented by 76).

25 211) The thrombopoietin receptor activators represented by 77).

212) The thrombopoietin receptor activators represented

by 78) .

213) The thrombopoietin receptor activators represented
by 79) .

214) The thrombopoietin receptor activators represented
5 by 80) .

215) The thrombopoietin receptor activators represented
by 81) .

216) The thrombopoietin receptor activators represented
by 82) .

10 217) The thrombopoietin receptor activators represented
by 83) .

218) The thrombopoietin receptor activators represented
by 84) .

219) The thrombopoietin receptor activators represented
15 by 85) .

220) The thrombopoietin receptor activators represented
by 86) .

221) The thrombopoietin receptor activators represented
by 87) .

20 222) The thrombopoietin receptor activators represented
by 88) .

223) The thrombopoietin receptor activators represented
by 89) .

224) The thrombopoietin receptor activators represented
25 by 90) .

225) The thrombopoietin receptor activators represented
by 91) .

226) The thrombopoietin receptor activators represented
by 92).

227) The thrombopoietin receptor activators represented
by 93).

5 228) The thrombopoietin receptor activators represented
by 94).

229) The thrombopoietin receptor activators represented
by 95).

10 230) The thrombopoietin receptor activators represented
by 96).

231) The thrombopoietin receptor activators represented
by 97).

232) The thrombopoietin receptor activators represented
by 98).

15 233) The thrombopoietin receptor activators represented
by 99).

234) The thrombopoietin receptor activators represented
by 100).

20 235) The thrombopoietin receptor activators represented
by 101).

236) The thrombopoietin receptor activators represented
by 102).

237) The thrombopoietin receptor activators represented
by 103).

25 238) The thrombopoietin receptor activators represented
by 104).

239) The thrombopoietin receptor activators represented

by 105) .

240) The thrombopoietin receptor activators represented
by 106) .

241) The thrombopoietin receptor activators represented
5 by 107) .

242) The thrombopoietin receptor activators represented
by 108) .

243) The thrombopoietin receptor activators represented
by 109) .

10 244) The thrombopoietin receptor activators represented
by 110) .

245) The thrombopoietin receptor activators represented
by 111) .

246) The thrombopoietin receptor activators represented
15 by 112) .

247) The thrombopoietin receptor activators represented
by 113) .

248) The thrombopoietin receptor activators represented
by 114) .

20 249) The thrombopoietin receptor activators represented
by 115) .

250) The thrombopoietin receptor activators represented
by 116) .

251) The thrombopoietin receptor activators represented
25 by 117) .

252) The thrombopoietin receptor activators represented
by 118) .

253) The thrombopoietin receptor activators represented
by 119).

254) The thrombopoietin receptor activators represented
by 120).

5 255) The thrombopoietin receptor activators represented
by 121).

256) The thrombopoietin receptor activators represented
by 122).

10 257) The thrombopoietin receptor activators represented
by 123).

258) The thrombopoietin receptor activators represented
by 124).

259) The thrombopoietin receptor activators represented
by 125).

15 260) The thrombopoietin receptor activators represented
by 126).

261) The thrombopoietin receptor activators represented
by 127).

20 262) The thrombopoietin receptor activators represented
by 128).

263) The thrombopoietin receptor activators represented
by 129).

264) The thrombopoietin receptor activators represented
by 130).

25 265) The thrombopoietin receptor activators represented
by 131).

266) The thrombopoietin receptor activators represented

by 132).

267) The thrombopoietin receptor activators represented by 133).

268) The thrombopoietin receptor activators represented
5 by 134).

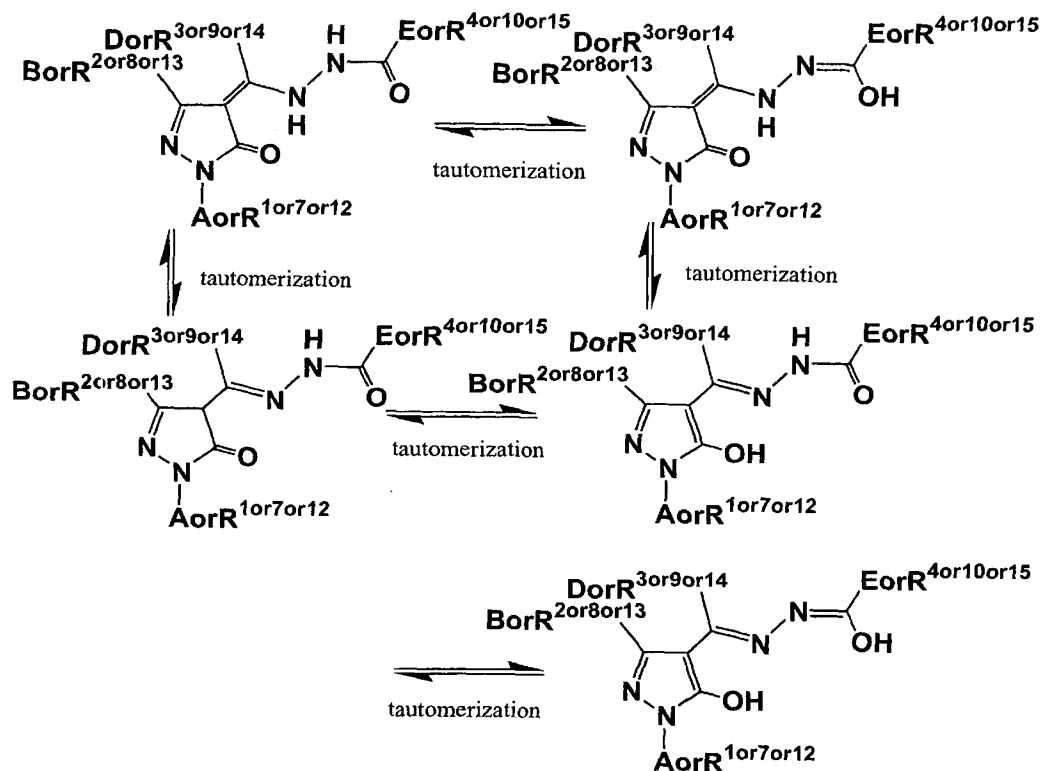
269) Preventive, therapeutic and improving agents for diseases against which activation of the thrombopoietin receptor is effective which contain the thrombopoietin receptor activators represented by any of 135) to 268) or
10 the formula (1), the formula (2), the formula (3) or the formula (4), tautomers, prodrugs or pharmaceutically acceptable salts of the activators or solvates thereof, as an active ingredient.

270) Platelet increasing agents containing the
15 thrombopoietin receptor activators represented by any of 135) to 268) or the formula (1), the formula (2), the formula (3) or the formula (4), tautomers, prodrugs or pharmaceutically acceptable salts of the activators or solvates thereof, as an active ingredient.

20 The compounds of the present invention represented by the formula (1), the formula (2), the formula (3) or the formula (4) may be present in the form of pyrazoles which undergo the following tautomerizations, mixtures or mixtures of isomers thereof. When the compounds of the
25 present invention have optical isomers, diastereomers or geometric isomers, the compounds of the present invention may be in the form of mixtures thereof or in the resolved

form.

【Ka 15】



The compounds of the present invention represented by
 5 the formula (1), the formula (2), the formula (3) or the
 formula (4) or pharmaceutically acceptable salts thereof
 may be in the form of arbitrary crystals or arbitrary
 hydrates. The present invention covers these crystals,
 hydrates and mixtures. They may be in the form of
 10 optional solvates with organic solvents such as acetone,
 ethanol and tetrahydrofuran, and the present invention
 covers any of these forms.

The compounds of the present invention represented by
 the formula (1), the formula (2), the formula (3) or the

formula (4) may be converted to pharmaceutically acceptable salts or may be liberated from the resulting salts, if necessary. The pharmaceutically acceptable salts of the present invention may be, for example, salts
5 with alkali metals (such as lithium, sodium and potassium), alkaline earth metals (such as magnesium and calcium), ammonium, organic bases and amino acids. They may be salts with inorganic acids (such as hydrochloric acid, hydrobromic acid, phosphoric acid and sulfuric
10 acid) and organic acids (such as acetic acid, citric acid, maleic acid, fumaric acid, benzenesulfonic acid and p-toluenesulfonic acid). They may also be complexes with transition metals (such as copper and zinc).

The compounds which serve as prodrugs are derivatives
15 of the present invention having chemically or metabolically degradable groups which give pharmacologically active compounds of the present invention upon solvolysis or under physiological conditions in vivo. Methods for selecting or producing
20 appropriate prodrugs are disclosed, for example, in Design of Prodrug (Elsevier, Amsterdam 1985). In the present invention, when the compound has a hydroxyl group, acyloxy derivatives obtained by reacting the compound with appropriate acyl halides or appropriate acid
25 anhydrides may, for example, be mentioned as a prodrug. Acyloxys particularly preferred as prodrugs include $-OCOC_2H_5$, $-OCO(t-Bu)$, $-OCOC_{15}H_{31}$, $-OCO(m-CO_2Na-Ph)$,

-OCOCH₂CH₂CO₂Na, -OCOCH(NH₂)CH₃, -OCOCH₂N(CH₃)₂ and the like. When the compound of the present invention has an amino group, amide derivatives obtained by reacting the compound having an amino group with appropriate acid
5 halides or appropriate mixed acid anhydrides may, for example, be mentioned as prodrugs. Amides particularly preferred as prodrugs include -NHCO(CH₂)₂₀OCH₃, -NHCOCH(NH₂)CH₃ and the like. When the compound of the present invention has a carboxyl group, carboxylic acid
10 esters with aliphatic alcohols or carboxylic acid esters obtained by the reaction of an alcoholic free hydroxyl group of 1,2- or 1,3-diglycerides may, for example, be mentioned as prodrugs. Particularly preferred prodrugs are methyl esters and ethyl esters.

15 The preventive, therapeutic and improving agents for diseases against which activation of the thrombopoietin receptor is effective or platelet increasing agents which contain the thrombopoietin receptor activators, tautomers, prodrugs or pharmaceutically acceptable salts of the
20 activators or solvates thereof as an active ingredient may usually be administered as oral medicines such as tablets, capsules, powder, granules, pills and syrup, as rectal medicines, percutaneous medicines or injections. The agents of the present invention may be administered
25 as a single therapeutic agent or as a mixture with other therapeutic agents. Though they may be administered as they are, they are usually administered in the form of

medical compositions. These pharmaceutical preparations can be obtained by adding pharmacologically and pharmaceutically acceptable additives by conventional methods. Namely, for oral medicines, ordinary excipients, lubricants, binders, disintegrants, humectants, plasticizers and coating agents may be used. Oral liquid preparations may be in the form of aqueous or oily suspensions, solutions, emulsions, syrups or elixirs or may be supplied as dry syrups to be mixed with water or other appropriate solvents before use. Such liquid preparations may contain ordinary additives such as suspending agents, perfumes, diluents and emulsifiers. In the case of rectal administration, they may be administered as suppositories. Suppositories may use an appropriate substance such as cacao butter, laurin tallow, Macrogol, glycerogelatin, Witepsol, sodium stearate and mixtures thereof as the base and may contain an emulsifier, a suspending agent, a preservative and the like. For injections, a solvent or a solubilizing agent such as distilled water for injection, physiological saline, 5% glucose solution and propylene glycol and pharmaceutical components such as a pH regulator, an isotonizing agent and a stabilizer may be used to form aqueous dosage forms or dosage forms which need dissolution before use.

The dose of the agents of the present invention for administration to human is usually about from 0.1 to 1000

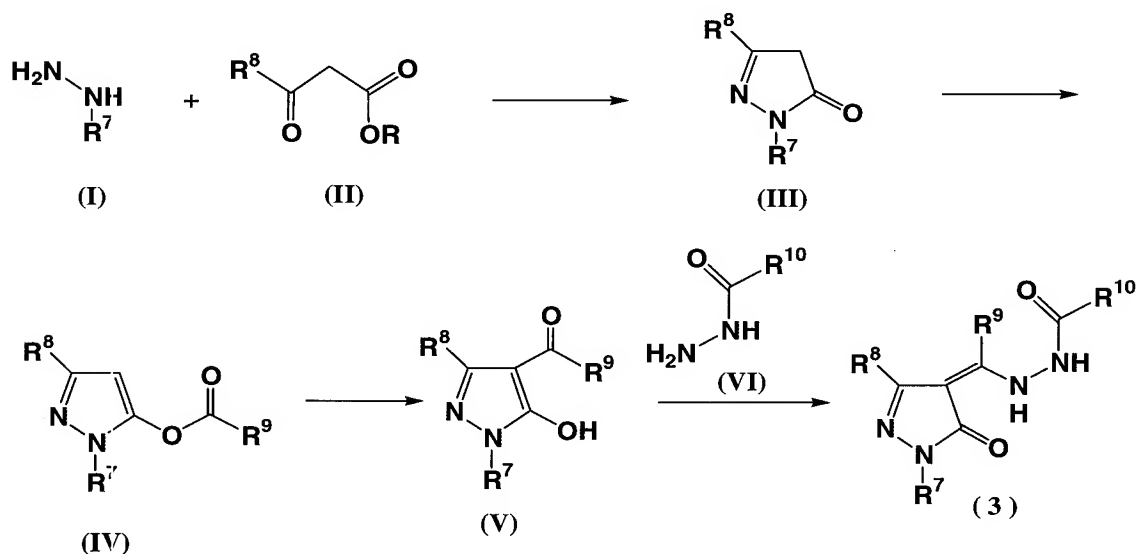
mg/human/day in the case of oral drugs or rectal administration and about from 0.05 mg to 500 mg/human/day in the case of injections, though it depends on the age and conditions of the patient. The above-mentioned
5 ranges are mere examples, and the dose should be determined from the conditions of the patient.

The present invention is used when the use of compounds which have thrombopoietin receptor affinity and act as thrombopoietin receptor agonists are expected to
10 improve pathological conditions. For example, hematological disorders accompanied by abnormal platelet count may be mentioned. Specifically, it is effective for therapy or prevention of human and mammalian diseases caused by abnormal megakaryopoiesis, especially those
15 accompanied by thrombocytopenia. Examples of such diseases include thrombocytopenia accompanying chemotherapy or radiotherapy of cancer, thrombocytopenia caused by bone marrow transplantation, surgery and serious infection, or gastrointestinal bleeding, but such
20 diseases are not restricted to these mentioned. Typical thrombocytopenias such as aplastic anemia, idiopathic thrombocytopenic purpura, myelodysplastic syndrome and thrombopoietin deficiency are also targets of the agents of the present invention. The present invention may be
25 used as a peripheral stem cell mobilizer, a megakaryocytic leukemia cell differentiation inducer and a platelet increasing agent for platelet donors. In

addition, potential applications include therapeutic
 angiogenesis based on differentiation and proliferation
 of vascular endothelial cells and endothelial progenitor
 cells, prevention and therapy of arteriosclerosis,
 5 myocardial infarction, unstable angina, peripheral artery
 occlusive disease, but there is no restriction.

The pyrazolone compounds represented by the formula
 (1), the formula (2), the formula (3) or the formula (4)
 are prepared by the process illustrated below in
 10 reference to the pyrazolone compounds represented by the
 formula (3).

【Ka 16】



The pyrazolones (III) are obtained by known methods
 15 (Syn. Comm 20(20), 3213 (1990), Chem Ber 59, 320 (1926),
 Monatsh. Chem 89, 30 (1958)), for example, by reacting β -
 keto esters (II) with hydrazines (R^7NHNH_2 or salts
 thereof) in acetic acid with reflux. Acylation of them

with acyl halides (R^9COCl) or acid anhydrides ($(R^9CO)_2O$) to (IV) followed by Fries rearrangement in the presence of potassium carbonate in dioxane with heating gives 4-acyl-5-hydroxypyrazoles (V). 4-Formyl-5-hydroxypyrazole (V) ($R^9 = H$) are obtainable by reacting the pyrazolones (III) with $POCl_3$ -DMF. They are heated with hydrazides ($R^{10}CCNHNH_2$ (VI) or salts thereof) optionally in the presence of a catalyst in a solvent to give the desired products. Syntheses of hydrazides (VI) are disclosed in the following documents.

- 1) Synthetic Commun., 28, (7) pp.1223-1231 (1998)
- 2) J. Chem. Soc., 1225 (1948)
- 3) J. Chem. Soc., 2831 (1952)
- 4) WO03/7328
- 5) Nihon Kagaku Zasshi, 88(5), p.73 (1967)
- 6) Journal of Heterocyclic Chemistry, 28(17), 17 (1991)

The compounds of the present invention are usually obtained with high purity by recrystallization or washing with solvents because most of them have good crystallizability. However, if necessary, they may be purified by column chromatography, thin layer chromatography, high performance liquid chromatography (HPLC) or high performance liquid chromatography-mass spectrometry (LC-MS).

【Examples】

Now, the present invention will be described in further detail with reference to Examples. However, it

should be understood that the present invention is by no means restricted by these specific Examples.

In high performance liquid chromatography-mass spectrometry (LC-MS), the retention time was measured
5 under the following conditions.

Column: Waters XTerra MSC18 4.6×50 mm

Eluent: H₂O:CH₃CN = 85:15 → 15:85

Syntheses of the compounds of Reference Synthetic Examples followed Examples 2-5 (pages 12-14) of
10 WO01/34585.

SYNTHETIC EXAMPLE 1

Synthesis of 2,4-dihydroxybenzoic N'-(1-(3-methyl-5-oxo-1-(4-iodophenyl)-1,5-dihydro-pyrazol-4-ylidene)-ethyl)-hydrazide

15 1.03 g (3 mmol) of 1-(5-hydroxy-1-(4-iodophenyl)-3-methyl-1H-pyrazol-4-yl)-ethanone and 505 mg (3 mmol) of 2,4-dihydroxybenzoic hydrazide were dissolved in 50 ml of DMSO and heated at 85°C for 9 hours with stirring. After cooling and evaporation of the solvent, the crude product
20 was recrystallized from chloroform/ether to give 1.39 g of the desired product as a pale brown solid (yield 94%).

¹H-NMR (ppm in DMSO-d₆)

δ = 2.36 (s, 3H), 2.42 (s, 3H), 2.54 (s, 3H), 6.36 (t, 1H, J = 2 Hz), 6.40 (d, 1H, J = 2 Hz), 7.68-7.76 (m, 3H), 7.86
25 (d, 2H, J = 9 Hz)

LC/MS

M⁺ = 492.27 (2.88 min)

SYNTHETIC EXAMPLE 2

Synthesis of 3,5-dihydroxybenzoic N'-(1-(1-(4-tert-butylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

5 From 1-(1-(4-tert-butylphenyl)-5-hydroxy-3-methyl-1H-pyrazol-4-yl)-ethanone and 3,5-dihydroxybenzoic hydrazide, 40.1 mg of the desired product was obtained in the same manner as in Synthetic Example 1 as a yellow solid (yield 40%).

10 ¹H-NMR (ppm in DMSO-d₆)

δ = 1.29 (s, 9H), 2.36 (s, 3H), 2.41 (s, 3H), 6.45 (s, 1H), 6.76 (s, 2H), 7.41 (d, 2H, J = 8.8 Hz), 7.89 (d, 2H, J = 8.8 Hz), 9.65 (s, 2H), 11.08 (s, 1H).

LC/MS

15 M⁺ = 422 (2.19 min).

SYNTHETIC EXAMPLE 3

Synthesis of 3,5-dihydroxybenzoic N'-(1-(1-(3,4-dimethylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

20 From 1-(1-(3,4-dimethylphenyl)-5-hydroxy-3-methyl-1H-pyrazol-4-yl)-ethanone and 3,5-dihydroxybenzoic hydrazide, 57.0 mg of the desired product was obtained in the same manner as in Synthetic Example 1 as a pale red solid (yield 73%).

25 ¹H-NMR (ppm in DMSO-d₆)

δ = 2.21 (s, 3H), 2.24 (s, 3H), 2.35 (s, 3H), 2.41 (s, 3H), 6.45 (s, 1H), 6.75 (s, 1H), 6.76 (s, 1H), 7.14 (d, 1H,

$J = 8.3$ Hz), 7.70 (dd, 1H, $J = 1.9, 8.3$ Hz), 7.77 (d, 1H, $J = 1.9$ Hz), 9.66 (s, 2H), 11.09 (s, 1H).

LC/MS

$M^+ = 394$ (1.82 min).

5 SYNTHETIC EXAMPLE 4

Synthesis of 4-methoxycarbonyl-benzoic N'-(1-(1-(4-tert-butylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

1) Synthesis of 4-methoxycarbonylbenzhydrazide

10 The known procedure disclosed in the literature (Synthetic Communications, 28(7), 1223-1231, (1998)) was followed using monomethyl terephthalate and tetramethylfluoroformamidinium hexafluorophosphate to give 1.36 g of a colorless solid (yield 70%).

15 $^1\text{H-NMR}$ (ppm in DMSO- d_6)

$\delta = 3.86$ (s, 3H), 4.56 (s, 2H), 7.93 (d, 2H, $J = 8.3$ Hz), 8.02 (d, 2H, $J = 8.3$ Hz), 9.96 (bs, 1H).

2) Synthesis of 4-methoxycarbonylbenzoic N'-(1-(1-(4-tert-butylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

20 30.5 mg (0.11 mmol) of 1-(1-(4-tert-butylphenyl)-5-hydroxy-3-methyl-1H-pyrazol-4-yl)-ethanone and 23.1 mg (0.11 mmol) of 4-methoxycarbonylbenzhydrazide were dissolved in 3.0 ml of DMF and stirred at 100°C for 3 hours. After cooling and evaporation of the solvent, the crude product was recrystallized from ethyl acetate/n-hexane to give 32.9 mg of the desired product as a yellow

solid (yield 66%).

¹H-NMR (ppm in DMSO-d₆)

δ = 1.29 (s, 9H), 2.37 (s, 3H), 2.46 (s, 3H), 3.90 (s,
3H), 7.41 (d, 2H, J = 8.7 Hz), 7.89 (d, 2H, J = 8.7 Hz),
5 8.05 (d, 2H, J = 8.4 Hz), 8.12 (d, 2H, J = 8.4 Hz).

LC/MS

M⁺ = 448 (2.64 min).

SYNTHETIC EXAMPLE 5

Synthesis of 4-carboxybenzoic N'-(1-(1-(4-tert-
10 butylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-
ylidene)-ethyl)-hydrazide

To 23.2 mg (0.05 mmol) of the 4-
methoxycarbonylbenzoic N'-(1-(1-(4-tert-butylphenyl)-3-
methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-
15 hydrazide synthesized in Synthetic Example 4 in 2.0 ml of
methanol, 255 μl (0.255 mmol) of 1M aqueous sodium
hydroxide was added at room temperature, and the mixture
was heated at from 60°C to 80°C for 3.5 hours. After it
was cooled to room temperature, 255 μl (0.255 mmol) of 1M
20 hydrochloric acid was added, and the precipitated solid
was collected by filtration to obtain 13.9 mg of the
desired product as a pale brown solid (yield 61%).

¹H-NMR (ppm in DMSO-d₆)

δ = 1.29 (s, 9H), 2.37 (s, 3H), 2.45 (s, 3H), 7.41 (d, 2H,
25 J = 8.7 Hz), 7.89 (d, 2H, J = 8.7 Hz), 8.03 (d, 2H, J =
8.3 Hz), 8.09 (d, 2H, J = 8.3 Hz), 11.44 (s, 1H).

LC/MS

M⁺ = 434 (2.38 min).

Synthetic EXAMPLE 6

Synthesis of 4-methoxycarbonylbenzoic N'-(1-(1-(3,4-dimethylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

From 1-(1-(3,4-dimethylphenyl)-5-hydroxy-3-methyl-1H-pyrazol-4-yl)-ethanone and 4-methoxycarbonylbenzhydrazide, 53.0 mg of the desired product was obtained in the same manner as in Synthetic Example 4 as a pale yellow solid (yield 64%).

¹H-NMR (ppm in DMSO-d₆)

δ = 2.21 (s, 3H), 2.25 (s, 3H), 2.36 (s, 3H), 2.45 (s, 3H), 3.89 (s, 3H), 7.14 (d, 1H, J = 8.5 Hz), 7.71 (dd, 1H, J = 1.9, 8.5 Hz), 7.77 (d, 1H, J = 1.9 Hz), 8.05 (d, 2H, J = 8.5 Hz), 8.12 (d, 2H, J = 8.5 Hz).

LC/MS

M⁺ = 420 (2.34 min).

SYNTHETIC EXAMPLE 7

Synthesis of 4-carboxybenzoic N'-(1-(1-(3,4-dimethylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

From the 4-methoxycarbonylbenzoic N'-(1-(1-(3,4-dimethylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide synthesized in Synthetic Example 6, 21.5 mg of the desired product was obtained in the same manner as in Synthetic Example 5 as a pale yellow solid (yield 71%).

¹H-NMR (ppm in DMSO-d₆)

δ = 2.21 (s, 3H), 2.25 (s, 3H), 2.36 (s, 3H), 2.45 (s,
3H), 7.14 (d, 1H, J = 8.3 Hz), 7.70 (dd, 1H, J = 1.9, 8.3
Hz), 7.77 (d, 1H, J = 1.9 Hz), 8.03 (d, 2H, J = 8.3 Hz),
5 8.10 (d, 2H, J = 8.3 Hz), 11.45 (s, 1H).

LC/MS

M⁺ = 406 (2.03 min).

SYNTHETIC EXAMPLE 8

Synthesis of 4-methoxycarbonylbenzoic N'-(1-(3-methyl-5-
10 oxo-1-(3-trifluoromethylphenyl)-1,5-dihydropyrazol-4-
ylidene)-ethyl)-hydrazide

From 1-(5-hydroxy-3-methyl-1-(3-
trifluoromethylphenyl)-1H-pyrazol-4-yl)-ethanone and 4-
methoxycarbonylbenzhydrazide, 59.9 mg of the desired
15 product was obtained in the same manner as in Synthetic
Example 4 as a yellow solid (yield 65%).

¹H-NMR (ppm in DMSO-d₆)

δ = 2.40 (s, 3H), 2.51 (s, 3H), 3.91 (s, 3H), 7.49 (d, 1H,
J = 7.4 Hz), 7.66 (dd, 1H, J = 8.0, 8.3 Hz), 8.06 (d, 2H,
20 J = 8.3 Hz), 8.13 (d, 2H, J = 8.3 Hz), 8.29 (d, 1H, J =
8.0 Hz), 8.45 (s, 1H), 11.55 (bs, 1H), 12.47 (bs, 1H).

LC/MS

M⁺ = 460.41 (2.69 min).

SYNTHETIC EXAMPLE 9

Synthesis of 4-carboxybenzoic N'-(1-(3-methyl-5-oxo-1-(3-
25 trifluoromethylphenyl)-1,5-dihydropyrazol-4-ylidene)-
ethyl)-hydrazide

From the 4-methoxycarbonylbenzoic N'-(1-(3-methyl-5-oxo-1-(3-trifluoromethylphenyl)-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide synthesized in Synthetic Example 8, 26.5 mg of the desired product was obtained in the same manner as in Synthetic Example 5 as a pale yellow solid (yield 78%).

¹H-NMR (ppm in DMSO-d₆)

δ = 2.41 (s, 3H), 2.51 (s, 3H), 7.49 (d, 1H, J = 8.0 Hz), 7.66 (dd, 1H, J = 8.0 Hz), 8.03 (d, 2H, J = 8.3 Hz), 8.10 (d, 2H, J = 8.3 Hz), 8.29 (d, 1H, J = 8.0 Hz), 8.45 (s, 1H), 11.52 (bs, 1H), 12.46 (bs, 1H).

LC/MS

M⁺ = 446.38 (2.29 min).

SYNTHETIC EXAMPLE 10

Synthesis of 4-methoxycarbonylbenzoic N'-(1-(3-methyl-5-oxo-1-(4-trifluoromethylphenyl)-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

From 1-(5-hydroxy-3-methyl-1-(4-trifluoromethylphenyl)-1H-pyrazol-4-yl)-ethanone and 4-methoxycarbonylbenzhydrazide, 58.9 mg of the desired product was obtained in the same manner as in Synthetic Example 4 as a yellow solid (yield 65%).

¹H-NMR (ppm in DMSO-d₆)

δ = 2.40 (s, 3H), 2.51 (s, 3H), 3.91 (s, 3H), 7.77 (d, 2H, J = 8.5 Hz), 8.06 (d, 2H, J = 8.5 Hz), 8.13 (d, 2H, J = 8.5 Hz), 8.26 (d, 2H, J = 8.5 Hz), 11.56 (bs, 1H), 12.46 (bs, 1H).

LC/MS

M⁺ = 460.41 (2.62 min).

SYNTHETIC EXAMPLE 11

Synthesis of 4-carboxybenzoic N'-(1-(3-methyl-5-oxo-1-(4-trifluoromethylphenyl)-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

From the 4-methoxycarbonylbenzoic N'-(1-(3-methyl-5-oxo-1-(4-trifluoromethylphenyl)-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide synthesized in Synthetic Example 10, 18.6 mg of the desired product was obtained in the same manner as in Synthetic Example 5 as a pale yellow solid (yield 68%).

¹H-NMR (ppm in DMSO-d₆)

δ = 2.40 (s, 3H), 2.51 (s, 3H), 7.77 (d, 2H, J = 8.7 Hz), 8.03 (d, 2H, J = 8.2 Hz), 8.10 (d, 2H, J = 8.2 Hz), 8.23 (d, 2H, J = 8.7 Hz), 11.53 (bs, 1H), 12.45 (bs, 1H).

LC/MS

M⁺ = 446.38 (2.31 min).

SYNTHETIC EXAMPLE 12

Synthesis of 3-carboxybenzoic N'-(1-(1-(4-tert-butylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

1) Synthesis of 3-methoxycarbonylbenzhydrazide

The procedure in Synthetic Example 4 was followed using monomethyl isophthalate and tetramethylfluoroformamidinium hexafluorophosphate to give 244.6 mg of a yellow solid (yield > 99%).

¹H-NMR (ppm in DMSO-d₆)

δ = 3.89 (s, 3H), 4.61 (bs, 2H), 7.62 (dd, 1H, J = 8.0 Hz), 8.08 (dd, 2H, J = 1.8, 8.0 Hz), 8.42 (d, 1H, J = 1.8 Hz), 9.98 (bs, 1H).

5 LC/MS

M⁺ = 194 (0.51 min).

2) Synthesis of 3-methoxycarbonylbenzoic N'-(1-(1-(4-tert-butylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

10 From 1-(1-(4-tert-butylphenyl)-5-hydroxy-3-methyl-1H-pyrazol-4-yl)-ethanone and 3-methoxycarbonylbenzhydrazide, 64.6 mg of the desired product was obtained in the same manner as in Synthetic Example 4 as a yellow solid (yield 70%).

15 3) Synthesis of 3-carboxybenzoic N'-(1-(1-(4-tert-butylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

From the 3-methoxycarbonylbenzoic N'-(1-(1-(4-tert-butylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide synthesized in 2), 11.2 mg of the desired product was obtained in the same manner as in Synthetic Example 5 as a pale brown solid (yield 50%).

¹H-NMR (ppm in DMSO-d₆)

25 δ = 1.29 (s, 9H), 2.37 (s, 3H), 2.45 (s, 3H), 7.42 (d, 2H, J = 8.8 Hz), 7.70 (dd, 1H, J = 7.8 Hz), 7.89 (d, 2H, J = 8.8 Hz), 8.16 (d, 1H, J = 6.9 Hz), 8.51 (s, 1H), 11.46 (bs, 1H).

LC/MS

M^+ = 434.49 (2.37 min).

SYNTHETIC EXAMPLE 13

Synthesis of 3-carboxybenzoic N'-(1-(1-(3,4-dimethylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

1) Synthesis of 3-methoxycarbonylbenzoic N'-(1-(1-(3,4-dimethylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

From 1-(1-(3,4-dimethylphenyl)-5-hydroxy-3-methyl-1H-pyrazol-4-yl)-ethanone and 3-methoxycarbonylbenzhydrazide, 27.4 mg of the desired product was obtained in the same manner as in Synthetic Example 4 as a pale yellow solid (yield 35%).

$^1\text{H-NMR}$ (ppm in DMSO-d_6)

δ = 2.21 (s, 3H), 2.25 (s, 3H), 2.34 (s, 3H), 2.36 (s, 3H), 3.92 (s, 3H), 7.14 (d, 1H, J = 8.3 Hz), 7.70–7.77 (m, 3H), 8.20 (d, 2H, J = 8.0 Hz), 8.51 (s, 1H), 11.49 (s, 1H).

2) Synthesis of 3-carboxybenzoic N'-(1-(1-(3,4-dimethylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

From the 3-methoxycarbonylbenzoic N'-(1-(1-(3,4-dimethylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide synthesized in 1), 17.2 mg of the desired product was obtained in the same manner as in Synthetic Example 5 as a pale yellow solid (yield 68%).

$^1\text{H-NMR}$ (ppm in DMSO-d_6)

$\delta = 2.21$ (s, 3H), 2.25 (s, 3H), 2.36 (s, 3H), 2.45 (s, 3H), 7.14 (d, 1H, $J = 8.5$ Hz), $7.68-7.77$ (m, 3H), $8.15-8.20$ (m, 2H), 8.19 (d, 1H, $J = 7.2$ Hz), 8.50 (s, 1H).

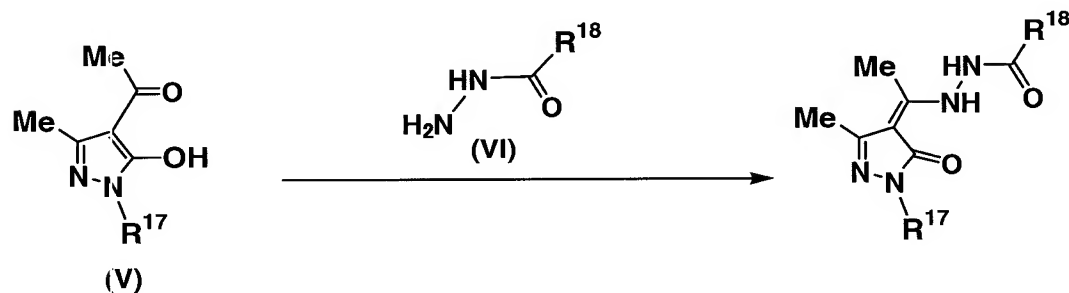
5 LC/MS

$M^+ = 406.43$ (2.03 min).

SYNTHETIC EXAMPLES 14 to 92

The structural formulae, yields, appearances, and molecular weights measured by LC/MS of the compounds synthesized by the following method based on Synthetic Example 1 are shown in Table 6.

【Ka 17】



A pyrazole derivative (V) and a benzoic hydrazide (VI) were dissolved in a solvent such as DMF, EtOH and DMSO in a molar ratio of 1:1 and heated at 80 to 100°C with stirring. The solvent was removed by evaporation, and the resulting crude product was dissolved in chloroform and recrystallized from a poor solvent or washed with chloroform to give the desired product.

Table 6

Syn- thet- ic Ex. No.	R ¹⁷	R ¹⁸	Yield	Appearance	Molec- ular weight
14	Ph	3-NO ₂ -Ph	37.6%	Yellow solid	379.38
15	4-t-Bu-Ph	3-NO ₂ -Ph	58.1%	Pale brown solid	435.48
16	Ph	2-OH-Ph	24.7%	Pale yellow solid	350.38
17	Ph	4-OH-Ph	65.1%	Pale pink solid	350.38
18	Ph	3-OH-2- Naphthyl	59.2%	Pale yellow solid	400.44
19	Ph	2,4-(OH) ₂ -Ph	41.1%	Pale yellow solid	366.38
20	Ph	3,4-(OH) ₂ -Ph	43.9%	Pale brown solid	366.38
21	Ph	2-NO ₂ -Ph	67.5%	Yellow solid	379.38
22	Ph	4-NO ₂ -Ph	53.4%	Yellow solid	379.38
23	4-t-Bu-Ph	2-OH-Ph	29.4%	Pale yellow solid	406.48
24	4-t-Bu-Ph	4-OH-Ph	24.1%	Pale brown solid	406.48
25	4-t-Bu-Ph	3-OH-2- Naphthyl	11.0%	Yellow solid	456.54
26	4-t-Bu-Ph	2,4-(OH) ₂ -Ph	27.5%	Pale yellow solid	422.48
27	4-t-Bu-Ph	3,4-(OH) ₂ -Ph	40.2%	Brown solid	422.48
28	4-t-Bu-Ph	2-NO ₂ -Ph	51.4%	Pale yellow solid	435.48

29	4-t-Bu-Ph	4-NO ₂ -Ph	49.9%	Yellow solid	435.48
30	4-CF ₃ -Ph	2-OH-Ph	48.5%	Yellow solid	418.37
31	4-CF ₃ -Ph	4-OH-Ph	60.0%	Pink solid	418.37
32	4-CF ₃ -Ph	3-OH-2-Naphthyl	8.2%	Pale yellow solid	468.43
33	4-CF ₃ -Ph	2,4-(OH) ₂ -Ph	3.1%	Brown solid	.434.37
34	4-CF ₃ -Ph	3,4-(OH) ₂ -Ph	73.2%	Pale pink solid	434.37
35	4-CF ₃ -Ph	2-NO ₂ -Ph	68.8%	Pale pink solid	447.37
36	4-CF ₃ -Ph	3-NO ₂ -Ph	64.2%	Pale yellow solid	447.37
37	4-CF ₃ -Ph	4-NO ₂ -Ph	60.1%	Pale yellow solid	447.37
38	4-I-Ph	2-OH-Ph	22.9%	Yellow solid	476.27
39	4-I-Ph	4-OH-Ph	36.6%	Pale brown solid	476.27
40	4-I-Ph	3-OH-2-Naphthyl	46.5%	Yellow solid	526.33
41	4-I-Ph	3,4-(OH) ₂ -Ph	52.5%	Pale pink solid	492.27
42	4-I-Ph	2-NO ₂ -Ph	43.3%	Pale pink solid	505.27
43	4-I-Ph	3-NO ₂ -Ph	51.4%	Yellow solid	505.27
44	4-I-Ph	4-NO ₂ -Ph	27.6%	Yellow solid	505.27
45	3-CF ₃ -Ph	2-OH-Ph	69.4%	Pale yellow solid	418.37
46	3-CF ₃ -Ph	4-OH-Ph	25.7%	Pale brown solid	418.37
47	3-CF ₃ -Ph	3-OH-2-Naphthyl	54.3%	Pale yellow solid	468.43

48	3-CF ₃ -Ph	2,4-(OH) ₂ -Ph	13.2%	Pale brown solid	434.37
49	3-CF ₃ -Ph	3,4-(OH) ₂ -Ph	57.3%	Pale pink solid	434.37
50	3-CF ₃ -Ph	2-NO ₂ -Ph	53.9%	Pink solid	447.37
51	3-CF ₃ -Ph	3-NO ₂ -Ph	57.4%	Pale yellow solid	447.37
52	3-CF ₃ -Ph	4-NO ₂ -Ph	32.2%	Pale yellow solid	447.37
53	3,4-Me ₂ -Ph	2-OH-Ph	52.2%	Pale yellow solid	378.43
54	3,4-Me ₂ -Ph	4-OH-Ph	66.2%	Pale pink solid	378.43
55	3,4-Me ₂ -Ph	3-OH-2- Naphthyl	65.9%	Pale yellow solid	428.49
56	3,4-Me ₂ -Ph	2,4-(OH) ₂ -Ph	43.0%	Pale yellow solid	394.43
57	3,4-Me ₂ -Ph	3,4-(OH) ₂ -Ph	40.4%	Pale yellow solid	394.43
58	3,4-Me ₂ -Ph	2-NO ₂ -Ph	67.9%	Pale yellow solid	407.43
59	3,4-Me ₂ -Ph	3-NO ₂ -Ph	50.8%	Pale yellow solid	407.43
60	3,4-Me ₂ -Ph	4-NO ₂ -Ph	67.1%	Pale brown solid	407.43
61	3,4-Cl ₂ -Ph	2-OH-Ph	45.6%	Pale yellow solid	419.27
62	3,4-Cl ₂ -Ph	4-OH-Ph	63.7%	Pale yellow solid	419.27
63	3,4-Cl ₂ -Ph	3-OH-2- Naphthyl	51.1%	Pale brown solid	469.33

64	3,4-Cl ₂ -Ph	2,4-(OH) ₂ -Ph	17.0%	Pale yellow solid	435.27
65	3,4-Cl ₂ -Ph	3,4-(OH) ₂ -Ph	66.1%	Pale pink solid	435.27
66	3,4-Cl ₂ -Ph	2-NO ₂ -Ph	67.4%	Pale yellow solid	448.27
67	3,4-Cl ₂ -Ph	3-NO ₂ -Ph	64.5%	Pale yellow solid	448.27
68	3,4-Cl ₂ -Ph	4-NO ₂ -Ph	51.1%	Brown solid	448.27
69	4-t-Bu-Ph	4-NH ₂ -Ph	74.8%	Pale brown solid	405.53
70	4-t-Bu-Ph	3-NH ₂ -Ph	48.7%	Pale brown solid	405.53
71	4-t-Bu-Ph	4-CF ₃ -Ph	69.1%	Pale yellow solid	458.49
72	4-t-Bu-Ph	4-t-Bu-Ph	77.9%	Pink solid	446.63
73	3,4-Me ₂ -Ph	4-NH ₂ -Ph	92.7%	Red solid	377.48
74	3,4-Me ₂ -Ph	3-NH ₂ -Ph	61.1%	Pale orange solid	377.48
75	3,4-Me ₂ -Ph	4-CF ₃ -Ph	67.7%	Pale orange solid	430.44
76	3,4-Me ₂ -Ph	4-t-Bu-Ph	66.8%	Pale pink solid	418.58
77	3,4-Cl ₂ -Ph	4-NH ₂ -Ph	51.2%	Orange solid	418.32
78	3,4-Cl ₂ -Ph	3-NH ₂ -Ph	69.7%	Pink solid	418.32
79	3,4-Cl ₂ -Ph	4-CF ₃ -Ph	69.6%	Pale orange solid	471.28
80	3,4-Cl ₂ -Ph	4-t-Bu-Ph	79.8%	Pale pink solid	459.42
81	4-t-Bu-Ph	3-OH-Ph	72.3%	Pale yellow solid	406.53

82	3,4-Me ₂ -Ph	3-OH-Ph	42.0%	Pale pink solid	378.48
83	3,4-Cl ₂ -Ph	3-OH-Ph	89.0%	Pink solid	419.32
84	3-NO ₂ -Ph	3-NO ₂ -Ph	58%	Brown solid	424.57
85	2-Py	3-NO ₂ -Ph	63%	Pale orange solid	380.36
86	3-NO ₂ -Ph	2,4-(OH) ₂ -Ph	43%	Brown solid	411.37
87	2-Py	2,4-(OH) ₂ -Ph	66%	Pale yellow solid	367.36
88	3-NO ₂ -Ph	4-t-Bu-Ph	25%	Brown solid	435.48
89	3-CF ₃ -Ph	3-NH ₂ -Ph	74%	Pale brown solid	417.38
90	3-CF ₃ -Ph	4-NH ₂ -Ph	82%	Pale orange solid	417.38
91	4-CF ₃ -Ph	3-NH ₂ -Ph	69%	Brown solid	417.38
92	4-CF ₃ -Ph	4-NH ₂ -Ph	72%	Pale pink solid	417.38

SYNTHETIC EXAMPLE 93

Synthesis of 2,4-dihydroxybenzoic N'-(1-(3,4-dimethylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene-methyl)-hydrazide

1) Synthesis of 1-(3,4-dimethylphenyl)-5-hydroxy-3-methyl-1H-pyrazole-4-carbaldehyde

1.86 g (9.16 mmol) of 1-(3,4-dimethylphenyl)-3-methyl-3-pyrazolin-5-one was dissolved in 3.6 ml of dry dimethylformamide, and 1.02 ml (11.0 mmol) of phosphorus oxychloride was added gradually under cooling with ice at

20°C or below. After the addition, the mixture was heated at 100°C for 2 hours, cooled to room temperature and poured into 30 ml of ice-cold water. Then, the mixture was washed with 10 ml of water and 10 ml of dimethylformamide. The mixed solution was stirred for 18 hours, and the precipitated solid was collected by filtration, washed with 20 ml of water and dried to obtain 1.03 g of the desired product as a pale brown solid (yield 49%).

10 ¹H-NMR (ppm in CDCl₃)

δ = 2.29 (s, 3H), 2.32 (s, 3H), 2.43 (s, 3H), 7.20 (d, 1H, J = 8 Hz), 7.48 (dd, 1H, J = 8 Hz, 2 Hz), 7.54 (d, 1H, J = 2 Hz), 9.60 (s, 1H)

2) Synthesis of 2,4-dihydroxybenzoic N'-[1-(3,4-dimethylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene-methyl]-hydrazide

46 mg (0.2 mmol) of the 1-(3,4-dimethylphenyl)-5-hydroxy-3-methyl-1H-pyrazole-4-carbaldehyde synthesized in 1) and 34 mg (0.20 mmol) of 2,4-dihydroxybenzoic hydrazide were stirred in 1 ml of ethanol at room temperature for 96 hours. The precipitated solid was collected by filtration and washed with 1 ml of ethanol, 1 ml of ether and 1 ml of methanol successively to obtain 53 mg of the desired product (yield 70%).

25 LC/MS

M⁺ = 380.40 (2.77 min)

SYNTHETIC EXAMPLE 94

Synthesis of 2,4-dihydroxybenzoic N'-(1-(1-(3,4-dimethylphenyl)-3-trifluoromethyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

5 1-(5-Hydroxy-3-methyl-1-(3-trifluoromethylphenyl)-1H-pyrazol-4-yl)-ethanone (0.173 mmol, 51.5 mg) and 2,4-dihydroxybenzoic hydrazide (0.173 mmol, 30.6 mg) were stirred in ethanol (5 ml) at 80°C for 19 hours. After the solvent was removed by evaporation, the residue was
10 dried with a vacuum pump and filtered with chloroform, and the filtrate was concentrated and resolved by silica gel thin layer chromatography (CHCl₃/MeOH = 10/1) to obtain 2,4-dihydroxybenzoic N'-(1-(1-(3,4-dimethylphenyl)-3-trifluoromethyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide as a pale
15 yellow solid (67 mg, yield 87%, purity 80.7%).

LC-MS 448.40 (M⁺)

SYNTHETIC EXAMPLE 95

Synthesis of 4-methoxycarbonylbenzoic N'-(1-(1-(3,4-dimethylphenyl)-3-trifluoromethyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

1-(5-Hydroxy-3-methyl-1-(3-trifluoromethylphenyl)-1H-pyrazol-4-yl)-ethanone (0.189 mmol, 56.5 mg) and 4-methoxycarbonylbenzhydrazide (0.189 mmol, 36.8 mg) were
25 stirred in DMF at 100°C for 2.2 hours and at 120°C for 17 hours. After the solvent was removed by evaporation, the residue was resolved by silica gel thin layer

chromatography (CHCl₃/MeOH = 10/1) to obtain 4-methoxycarbonylbenzoic N'-(1-(1-(3,4-dimethylphenyl)-3-trifluoromethyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide as a yellow solid (55.6 mg, 62%)

5 LC-MS 474.43 (M⁺)

SYNTHETIC EXAMPLE 96

Synthesis of 4-carboxybenzoic N'-(1-(1-(3,4-dimethylphenyl)-3-trifluoromethyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

10 4-Methoxycarbonylbenzoic N'-(1-(1-(3,4-dimethylphenyl)-3-trifluoromethyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide (0.107 mmol, 50.7 mg) was dissolved in methanol (2 ml) and stirred with 1M aqueous sodium hydroxide (0.534 mmol, 0.534 ml) at room temperature for 2 hours and at 60°C for 1.5 hours. Then, the reaction vessel was cooled to 0°C, and 1M hydrochloric acid (0.534 mmol, 0.534 ml) and water were added. The precipitated solid was collected by filtration with water and dried to obtain 4-carboxybenzoic N'-(1-(1-(3,4-dimethylphenyl)-3-trifluoromethyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide as a yellow solid (43.8 mg, 89%).

LC-MS 460.41 (M⁺)

SYNTHETIC EXAMPLE 97

25 Synthesis of 4-carboxy-benzoic N'-(1-(4-tert-butylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene-methyl)-hydrazide

1) Synthesis of 1-(4-tert-butylphenyl)-5-hydroxy-3-methyl-1H-pyrazole-4-carbaldehyde

1.89 g (9.33 mmol) of 1-(4-tert-butylphenyl)-5-hydroxy-3-methyl-1H-pyrazole was dissolved in 3.6 ml of dry dimethylformamide, and 1.05 ml (11.26 mmol) phosphorus oxychloride was added gradually at 20°C or below under cooling with ice. After the addition, the mixture was heated at 100°C for 3 hours, then cooled to room temperature and poured into 30 ml of ice-cold water. The mixed solution was stirred at room temperature for 18 hours, and the precipitated solid was collected by filtration, washed with 20 ml of water and dried to obtain 1.61 g of the above-identified desired product as a yellow solid (yield 70%).

¹H-NMR (ppm in DMSO- d₆)

δ = 1.30-1.33 (m, 9H), 2.34-2.44 (m, 3H), 7.48-7.62 (m, 4H), 9.62-9.90 (m, 1H).

2) Synthesis of 4-methoxycarbonyl-benzoic N'-(1-(4-tert-butylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene-methyl)-hydrazide

1.0712 g (4.21 mmol) of the 1-(4-tert-butylphenyl)-5-hydroxy-3-methyl-1H-pyrazole-4-carbaldehyde synthesized in 1) and 819.6 mg (4.22 mmol) of 4-methoxycarbonylbenzhydrazide were stirred in 10 ml of dimethylformamide at room temperature for 3 hours. After the solvent was removed by evaporation, the precipitated solid was washed with a small amount of methanol and

dried to obtain 765.9 mg of the above-identified desired product as a yellow solid (yield 42%).

¹H-NMR (ppm in DMSO- d₆)

δ = 1.30 (s, 9H), 2.19-2.21 (m, 3H), 3.90 (s, 3H), 7.33
5 (s, 1H), 7.40-7.46 (m, 2H), 7.81-7.89 (m, 2H), 8.01-8.17 (m, 4H).

3) Synthesis of 4-carboxy-benzoic N'-(1-(4-tert-butylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene-methyl)-hydrazide

10 59.4 mg (0.14 mmol) of the 4-methoxycarbonyl-benzoic N'-(1-(4-tert-butylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene-methyl)-hydrazide synthesized in 2) was dissolved in 5.0 ml of methanol and stirred with 0.68 ml (0.68 mmol) of 1M aqueous sodium hydroxide at
15 room temperature for 6 hours and then at 60°C for 3 hours. After the stirring, 0.68 ml (0.68 mmol) of hydrochloric acid was added, and the precipitated solid was collected by filtration and dried to obtain 33.3 mg of the above-identified desired product as a yellow solid (yield 58%).

20 ¹H-NMR (ppm in DMSO- d₆)

δ = 1.30 (s, 9H), 2.19-2.21 (m, 3H), 7.33 (s, 1H), 7.40-7.46 (m, 2H), 7.80-7.89 (m, 2H), 7.99-8.14 (m, 4H).

LC/MS

M⁺ = 420.46 (2.39 min)

25 SYNTHETIC EXAMPLE 98

Synthesis of 5-methoxycarbonyl-2-thiophenecarboxylic acid N'-(1-(1-(4-tert-butylphenyl)-3-methyl-5-oxo-1,5-

dihydropyrazol-4-ylidene)-ethyl)-hydrazide

1) Synthesis of 5-methoxycarbonyl-2-thiophenecarboxylic acid

1.72 g (10 mmol) of thiophene-2,5-dicarboxylic acid
5 and 3.18 g (30 mmol) of sodium carbonate suspended in 25 mL of DMF were stirred with 623 μ L of methyl iodide at room temperature overnight. The sodium salt of the desired product was extracted with water, and 12M of hydrochloric acid was added to the combined aqueous layer.
10 The desired product was extracted with ethyl acetate, and the combined organic layer was washed with saturated aqueous ammonium chloride and dried over anhydrous magnesium sulfate. The desired product was purified by silica gel column chromatography to give 0.49 g of a
15 colorless solid (yield 28%).

$^1\text{H-NMR}$ (ppm in CDCl_3)

δ = 3.93 (s, 3H), 7.77 (d, 1H, J = 4.2 Hz), 7.83 (d, 1H, J = 4.2 Hz).

LC/MS

20 M^+ = 186 (0.92 min)

2) Synthesis of 5-methoxycarbonyl-2-thiophenecarboxylic acid hydrazide

The known procedure disclosed in the literature (J. Heterocyclic Chem., 28, 17, (1991).) was followed using
25 5-methoxycarbonyl-2-thiophenecarboxylic acid, thionyl chloride and hydrazine monohydrate to give 144 mg of a white solid (yield 72%).

¹H-NMR (ppm in DMSO- d₆)

δ = 3.84 (s, 3H), 4.57 (brs, 2H), 7.72 (d, 1H, J = 4.2 Hz), 7.79 (d, 1H, J = 4.2 Hz), 10.06 (brs, 1H).

LC/MS

5 M⁺ = 200 (3.09 min)

3) Synthesis of 5-methoxycarbonyl-2-thiophenecarboxylic acid N'-(1-(1-(4-tert-butylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

54.5 mg (0.20 mmol) of 1-(1-(4-tert-butylphenyl)-5-hydroxy-3-methyl-1H-pyrazol-4-yl)-ethanone and 40.0 mg (0.20 mmol) of 5-methoxycarbonyl-2-thiophenecarboxylic acid hydrazide were dissolved in 2.0 mL of DMF and stirred at 110°C for 12 hours. After cooling, the solvent was removed by evaporation, and the crude product was washed with ethyl acetate and collected by filtration to obtain 32.0 mg of the desired product as a yellow solid (yield 35%).

¹H-NMR (ppm in DMSO- d₆)

δ = 1.29 (s, 9H), 2.36 (s, 3H), 2.43 (s, 3H), 3.87 (s, 3H), 7.41 (d, 2H, J = 9.0 Hz), 7.87-7.90 (m, 4H).

LC/MS

M⁺ = 454.54 (4.46 min)

SYNTHETIC EXAMPLE 99

Synthesis of 5-carboxy-2-thiophenecarboxylic acid N'-(1-(1-(4-tert-butylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

14.9 mg (0.033 mmol) of 5-methoxycarbonyl-2-

thiophenecarboxylic acid N'-(1-(1-(4-tert-butylphenyl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide in 1.5 mL of methanol was stirred with 164 μ L (0.164 mmol) of 1M aqueous sodium hydroxide at room temperature for 17 hours. After the stirring, 164 μ L (0.164 mmol) of 1M hydrochloric acid was added, and the precipitated solid was collected by filtration to obtain 6.8 mg of the desired product as a pale yellow solid (yield 47%).

¹H-NMR (ppm in DMSO- d₆)

δ = 1.29 (s, 9H), 2.36 (s, 3H), 2.43 (s, 3H), 7.41 (d, 2H, J = 9.0 Hz), 7.80 (d, 1H, J = 3.9 Hz), 7.87-7.90 (m, 3H).

LC/MS

M⁺ = 440.52 (4.23 min)

SYNTHETIC EXAMPLE 100

Synthesis of 4-carboxy-benzoic N'-(1-(1-(quinolin-2-yl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide

1) Synthesis of 4-methoxycarbonyl-benzoic N'-(1-(1-(quinolin-2-yl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide (compound)

2.0 mL of an isopropyl alcohol solution of 28.7 mg (0.11 mmol) of 1-(1-(quinolin-2-yl)-5-hydroxy-3-methyl-1H-pyrazol-4-yl)-ethanone, 20.8 mg (0.11 mmol) of 4-methoxycarbonylbenzhydrazide and 6.1 mg (0.03 mmol) of p-toluenesulfonic acid monohydrate was refluxed with heating for 48 hours. After cooling, the precipitate was

collected by filtration and washed with methanol and acetonitrile to obtain 14.9 mg of the desired product as a purple solid (yield 31%).

¹H-NMR (ppm in DMSO- d₆)

5 δ = 2.54 (s, 3H), 3.91 (s, 3H), 7.58-7.63 (m, 1H), 7.80-7.85 (m, 1H), 8.01-8.15 (m, 6H), 8.46 (d, 1H, J = 6.3 Hz), 8.58 (d, 1H, J = 6.3 Hz).

LC/MS

M⁺ = 443.45 (3.21 min)

10 2) Synthesis of 4-carboxy-benzoic N'-(1-(1-quinolin-2-yl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide (compound)

1.5 mL of a methanol solution of 14.9 mg (0.034 mmol) of the 4-methoxycarbonyl-benzoic N'-(1-(1-(quinolin-2-yl)-3-methyl-5-oxo-1,5-dihydropyrazol-4-ylidene)-ethyl)-hydrazide synthesized in 1) was stirred with 168 μL (0.168 mmol) of 1M aqueous sodium hydroxide at 50°C for 12 hours. After the stirring, 168 μL (0.168 mmol) of 1M hydrochloric acid was added, and the precipitated solid
15
20 was collected by filtration to obtain 4.9 mg of the desired product as a dark yellow solid (yield 34%).

¹H-NMR (ppm in DMSO- d₆)

δ = 2.44 (s, 3H), 7.52-7.56 (m, 1H), 7.75 (t, 1H, J = 7.5 Hz), 7.94 (d, 1H, J = 4.5 Hz), 7.96 (d, 1H, J = 4.2 Hz),
25 8.05 (d, 2H, J = 8.7 Hz), 8.10 (d, 2H, J = 8.4 Hz), 8.33 (d, 1H, J = 9.6 Hz), 8.42 (d, 1H, J = 9.0 Hz).

LC/MS

M⁺ = 429.43 (3.21 min)

SYNTHETIC EXAMPLE 101

Synthesis of methyl 4-[(2-{1-[1-(6-chloro-3-pyridazinyl)-
5-hydroxy-3-methyl-1H-pyrazol-4-
5 yl]ethylidene}hydrazino)carbonyl]benzoate

0.2 mmol of 1-[1-(6-chloro-3-pyridazinyl)-5-hydroxy-
3-methyl-1H-pyrazol-4-yl]ethanone and 0.2 mmol of 4-
methoxycarbonylbenzhydrazide were dissolved in 2 ml of
DMSO and heated at 100°C for 8 hours with stirring.

10 After the solvent was removed by evaporation, the crude
product was dissolved in chloroform and recrystallized
from ether to obtain 55 mg of the desired product, methyl
4-[(2-{1-[1-(6-chloro-3-pyridazinyl)-5-hydroxy-3-methyl-
1H-pyrazol-4-yl]ethylidene}hydrazino)carbonyl]benzoate
15 (yield 64%).

¹H-NMR (ppm in DMSO-d₆)

δ = 2.42 (s, 3H), 2.54 (s, 3H), 3.91 (s, 3H), 7.96 (d, 1H,
J = 9.3 Hz), 8.06 (d, 2H, J = 8.4 Hz), 8.13 (d, 2H, J =
8.4 Hz), 8.44 (d, 1H, J = 9.3 Hz).

20 LC/MS

M⁺ = 428.83 (2.88 min).

SYNTHETIC EXAMPLE 102

Synthesis of 4-{[2-(1-{5-hydroxy-3-methyl-1-[5-
(trifluoromethyl)-2-pyridinyl]-1H-pyrazol-4-
25 yl}ethylidene)hydrazino]carbonyl]benzoic acid

1) Synthesis of methyl 4-{[2-(1-{5-hydroxy-3-methyl-1-[5-
(trifluoromethyl)-2-pyridinyl]-1H-pyrazol-4-

yl}ethylidene)hydrazino]carbonyl}benzoate

0.2 mmol of 1-{5-hydroxy-3-methyl-1-[5-(trifluoromethyl)-2-pyridinyl]-1H-pyrazol-4-yl}ethanone and 0.2 mmol of 4-methoxycarbonylbenzhydrazide were

5 heated in 2 ml of DMF at 100°C for 9 hours with stirring. After the solvent was removed by evaporation, the resulting crude product was dissolved in chloroform and recrystallized from hexane to obtain 66 mg of the desired product, methyl 4-{[2-(1-{5-hydroxy-3-methyl-1-[5-(trifluoromethyl)-2-pyridinyl]-1H-pyrazol-4-yl}ethylidene)hydrazino]carbonyl}benzoate (yield 72%).

¹H-NMR (ppm in DMSO-d₆)

δ = 2.41 (s, 3H), 2.50 (s, 3H), 3.88 (s, 3H), 7.9-8.4 (m, 6H), 8.80 (s, 1H).

15 LC/MS

M⁺ = 461.39 (3.00 min).

2) Synthesis of 4-{[2-(1-{5-hydroxy-3-methyl-1-[5-(trifluoromethyl)-2-pyridinyl]-1H-pyrazol-4-yl}ethylidene)hydrazino]carbonyl}benzoic acid

20 50 mg of the methyl 4-{[2-(1-{5-hydroxy-3-methyl-1-[5-(trifluoromethyl)-2-pyridinyl]-1H-pyrazol-4-yl}ethylidene)hydrazino]carbonyl}benzoate synthesized in 1) was heated in 3 ml of methanol and 0.3 ml of 1M aqueous sodium hydroxide at 60°C for 8 hours with stirring. After it was cooled to room temperature, 0.3 ml of 1M hydrochloric acid was added to precipitate crystals, and crystals were collected by filtration and

dried to obtain 30 mg of the desired product, 4-{[2-(1-{5-hydroxy-3-methyl-1-[5-(trifluoromethyl)-2-pyridinyl]-1H-pyrazol-4-yl}ethylidene)hydrazino]carbonyl}benzoic acid as a pale brown solid (yield 62%).

5 ¹H-NMR (ppm in DMSO-d₆)

δ = 2.41 (3H, s), 2.50 (3H, s), 8.04 (d, 2H, J = 8.4 Hz), 8.10 (d, 2H, J = 8.4 Hz), 8.26 (dd, 1H, J = 9 Hz, J = 2.4 Hz), 8.35 (d, 1H, J = 9 Hz), 8.81 (brs, 1H), 11.6 (brs, 1H), 12.4 (brs, 1H)

10 LC/MS

M⁺ = 447.37 (2.65 min).

REFERENCE SYNTHETIC EXAMPLE 1 (EXAMPLE 4 OF WO01/34585)

Synthesis of 5-(4-carboxybenzylidene)-3-[(1-{3,4-dimethylphenyl}-5-hydroxy-3-methyl-1H-pyrazol-4-

15 ylmethylene)amino]-2-thioxothiazolidin-4-one

1) Synthesis of 1-(3,4-dimethylphenyl)-5-hydroxy-3-methyl-1H-pyrazole-4-carbaldehyde

1.86 g (9.16 mmol) of 1-(3,4-dimethylphenyl)-3-methyl-3-pyrazolin-5-one was dissolved in 3.6 ml of dry
20 dimethylformamide, and 1.02 ml (11.0 mmol) of phosphorus oxychloride was added gradually under cooling with ice at 20°C or below. After the addition, the mixture was heated at 100°C for 2 hours, then cooled to room temperature and poured into 30 ml of ice-cold water.
25 Then, it was washed with 10 ml of water and 10 ml of dimethylformamide. The mixed solution was stirred for 18 hours, and the precipitated solid was collected by

filtration, washed with 20 ml of water and dried to obtain 1.03 g of the above-identified desired product as a pale brown solid (yield 49%).

¹H-NMR (ppm in CDCl₃)

5 δ = 2.29 (s, 3H), 2.32 (s, 3H), 2.43 (s, 3H), 7.20 (d, 1H, J = 8 Hz), 7.48 (dd, 1H, J = 8 Hz, 2Hz), 7.54 (d, 1H, J = 2 Hz), 9.60 (s, 1H)

2) Synthesis of 5-(4-carboxybenzylidene)-3-[(1-{3,4-dimethylphenyl}-5-hydroxy-3-methyl-1H-pyrazol-4-ylmethylene)amino]-2-thioxothiazolidin-4-one

230 mg (1 mmol) of the 1-(3,4-dimethylphenyl)-5-hydroxy-3-methyl-1H-pyrazole-4-carbaldehyde synthesized in 1) and 148 mg (1 mmol) of 3-aminorhodanine were stirred in 10 ml of ethanol at room temperature for 96
15 hours. The resulting solid was collected by filtration, washed with ethanol and ether and dried to obtain 332 mg of a crude imine.

A liquid mixture of 160 mg (0.444 mmol) of the imine, 4 mg of piperidine, 66 mg of 4-formylbenzoic acid, 6 mg
20 of benzoic acid and 20 ml of toluene was refluxed in a reactor equipped with a Dean-Stark tube packed with molecular sieve for 7 hours with heating. After cooling, the precipitated solid was collected by filtration and washed with 3 ml of toluene and 3 ml of ether to obtain
25 23.3 mg of a yellow solid. It was washed with a liquid mixture of methanol and chloroform to obtain 16.5 mg of the desired product (yield 7.5%).

¹H-NMR (ppm in DMSO-d₆)

δ = 2.10-2.40 (s×3, 9H), 7.18(d, 1H, J = 8 Hz), 7.63 (d, 1H, J = 8 Hz), 7.67 (s, 1H), 7.84 (d, 2H, J = 8 Hz), 8.03 (d, 2H, J = 8 Hz), 8.10 (d, 2H, J = 8 Hz), 8.20 (s, 1H)

5 LC/MS

M⁺ = 493.0 (3.33 min)

REFERENCE SYNTHETIC EXAMPLE 2 (EXAMPLE 5 OF WO01/34585)

Synthesis of 5-(3-carboxybenzylidene)-3-[(1-{3,4-dimethylphenyl}-5-hydroxy-3-methyl-1H-pyrazol-4-ylmethylene)amino]-2-thioxothiazolidin-4-one

A liquid mixture of 160 mg (0.444 mmol) of the imine synthesized in 2) of Reference Synthetic Example 1, 4 mg of piperidine, 66 mg of 3-formylbenzoic acid, 6 mg of benzoic acid and 20 ml of toluene was refluxed in a reactor equipped with a Dean-Stark tube packed with molecular sieve for 7 hours with heating. After cooling, the precipitated solid was collected by filtration and washed with 3 ml of toluene and 3 ml of ether to obtain 38.5 mg of a yellow solid (yield 18%).

20 ¹H-NMR (ppm in DMSO-d₆)

δ = 2.00-2.30 (s×3, 9H), 7.18 (d, 1H, J = 8 Hz), 7.64 (d, 1H, J = 8 Hz), 7.68 (s, 1H), 7.73 (t, 1H, J = 8 Hz), 7.97 (d, 2H, J = 8 Hz), 8.06 (s, 1H), 8.08 (d, 1H, J = 8 Hz), 8.23 (d, 2H, J = 8 Hz)

25 LC/MS

M⁺ = 493.0 (3.32 min)

REFERENCE SYNTHETIC EXAMPLE 3 (EXAMPLE 2 OF WO01/34585)

Synthesis of 3-(3-carboxyphenyl)-1-[(1-(3,4-dimethylphenyl)-5-hydroxy-3-methyl-1H-pyrazol-4-ylmethylene)amino]-2-thioxoimidazolidin-4-one

- 5 1) Synthesis of 1-amino-3-(3-carboxyphenyl)-2-thioxoimidazolidin-4-one

179 mg (1 mmol) of 3-isothiocyanatobenzoic acid and 523 μ l (3 mmol) of diisopropylethylamine were stirred in 8 ml of dichloromethane and then with 155 mg (1 mmol) of ethyl hydrazinoacetate hydrochloride at room temperature for 96 hours. After the solvent was concentrated, the mixture was partitioned between ethyl acetate and 30% acetic acid. The aqueous layer was extracted with ethyl acetate again, and the organic layers were combined, washed with water and then with saturated aqueous sodium chloride, dried over magnesium sulfate and concentrated. The resulting solid was mixed with a 190:10:0.8 liquid mixture of ethyl acetate, methanol and acetic acid, and the insoluble was dried to obtain 55.7 mg of the desired product (yield 22%).

^1H -NMR (ppm in DMSO- d_6)

δ = 4.44 (s, 2H), 5.46 (s, 2H), 7.57 (dd, 1H, J = 8 Hz, J = 1.5 Hz), 7.63 (t, 1H, J = 8 Hz), 7.90 (s, 1H), 7.99 (d, 1H, J = 8 Hz)

25 LC/MS

M^+ = 251.30 (0.59 min).

2) Synthesis of 3-(3-carboxyphenyl)-1-[(1-(3,4-

dimethylphenyl)-5-hydroxy-3-methyl-1H-pyrazol-4-ylmethylene)amino]-2-thioxoimidazolidin-4-one

50 mg (0.2 mmol) of the 1-amino-3-(3-carboxyphenyl)-2-thioxoimidazolidin-4-one synthesized above in 1) and 55 mg (0.22 mmol) of the 1-(3,4-dimethylphenyl)-5-hydroxy-3-methyl-1H-pyrazole-4-carbaldehyde synthesized in 1) of Reference Synthetic Example 1 were stirred in a liquid mixture of 10 ml of ethanol and 5 ml of methanol at room temperature for 96 hours. The resulting insoluble was collected by filtration to obtain 73 mg of the desired product as a yellow solid (yield 72%).

¹H-NMR (ppm in DMSO-d₆)

δ = 2.24 (s, 3H), 2.27 (s, 3H), 2.38 (s, 3H), 4.74 (s, 2H), 7.21 (d, 1H, J = 8 Hz), 7.40-7.80 (m, 4H), 7.95 (s, 1H), 8.02 (d, 1H, J = 8 Hz), 8.14 (s, 1H)

LC/MS

M⁺ = 463.51 (2.77 min).

REFERENCE SYNTHETIC EXAMPLE 4 (EXAMPLE 3 OF WO01/34585)

Synthesis of 3-(4-carboxyphenyl)-1-[(1-(3,4-dimethylphenyl)-5-hydroxy-3-methyl-1H-pyrazol-4-ylmethylene)amino]-2-thioxoimidazolidin-4-one

1) Synthesis of 1-amino-3-(4-carboxyphenyl)-2-thioxoimidazolidin-4-one

179 mg (1 mmol) of 4-isothiocyanatobenzoic acid and 523 μl (3 mmol) of diisopropylethylamine were stirred in 8 ml of dichloromethane and then with 155 mg (1 mmol) of ethyl hydrazinoacetate hydrochloride at room temperature

for 96 hours. After the solvent was concentrated, the mixture was partitioned between ethyl acetate and 30% acetic acid. The aqueous layer was extracted with ethyl acetate again, and the organic layers were combined, washed with water and then with saturated aqueous sodium chloride, dried over magnesium sulfate and concentrated. The resulting solid was mixed with a 190:10:0.8 liquid mixture of ethyl acetate, methanol and acetic acid, and the insoluble was dried to obtain 132 mg of the desired product (yield 53%).

$^1\text{H-NMR}$ (ppm in DMSO-d_6)

δ = 4.46(s, 2H), 5.47 (s, 2H), 7.46 (d, 2H, J = 8 Hz), 8.04 (d, 2H, J = 8 Hz)

LC/MS

M^+ = 251.26 (0.95 min).

2) Synthesis of 3-(4-carboxyphenyl)-1-[(1-(3,4-dimethylphenyl)-5-hydroxy-3-methyl-1H-pyrazol-4-ylmethylene)amino]-2-thioxoimidazolidin-4-one

50 mg (0.2 mmol) of the 1-amino-3-(4-carboxyphenyl)-2-thioxoimidazolidin-4-one synthesized above in 1) and 55 mg (0.22 mmol) of the 1-(3,4-dimethylphenyl)-5-hydroxy-3-methyl-1H-pyrazole-4-carbaldehyde synthesized in 1) of Reference Synthetic Example 1 were stirred in a liquid mixture of 10 ml of ethanol and 5 ml of methanol at room temperature for 96 hours. The resulting insoluble was collected by filtration to obtain 87 mg of the desired product as a yellow solid (yield 85%).

^1H -NMR (ppm in DMSO- d_6)

δ = 2.24 (s, 3H), 2.27 (s, 3H), 2.50 (s, 3H), 4.75 (s, 2H), 7.21 (d, 1H, J = 8 Hz), 7.40-7.70 (m, 4H), 8.08 (d, 2H, J = 8.8 Hz), 8.14 (brs, 1H)

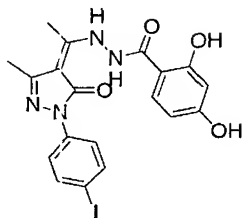
5 LC/MS

M^+ = 463.51 (2.76 min).

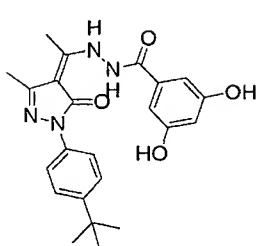
The structural formulae of the compounds obtained in the Synthetic Examples are as follows.

【Ka 18】

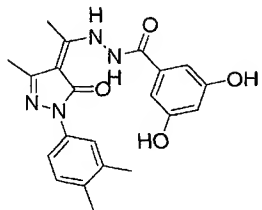
SYNTHETIC EX. 1



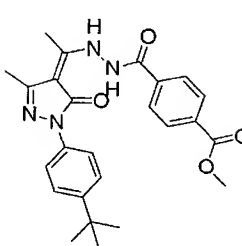
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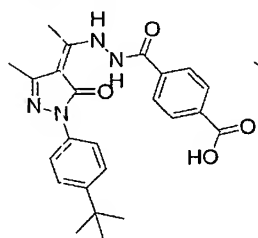
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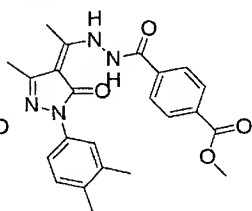
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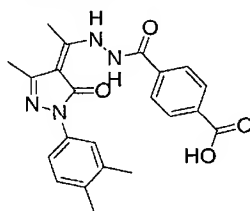
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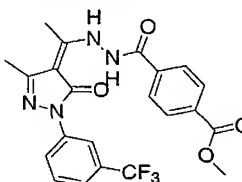
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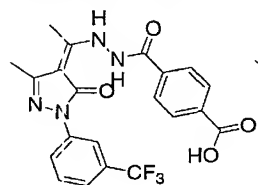
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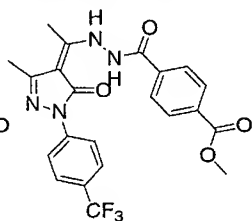
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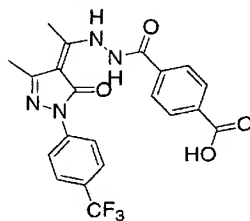
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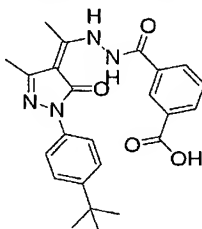
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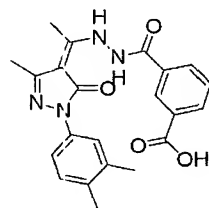
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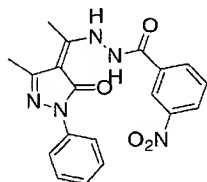
SYNTHETIC EX. 12



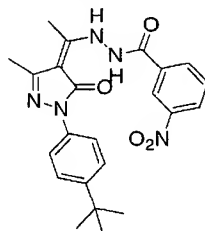
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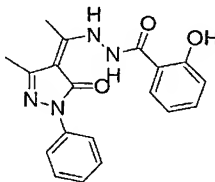
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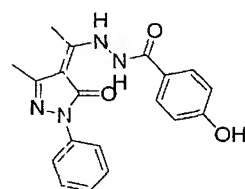
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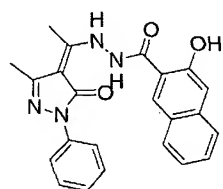
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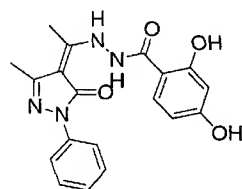
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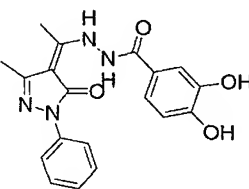
SYNTHETIC EX. 18



SYNTHETIC EX. 19

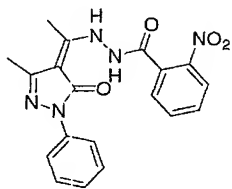


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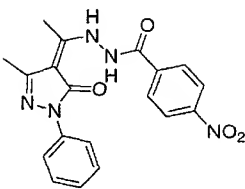


【Ka 19】

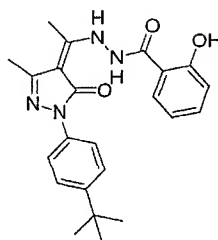
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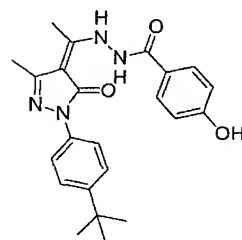
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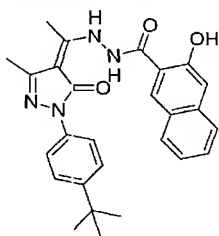
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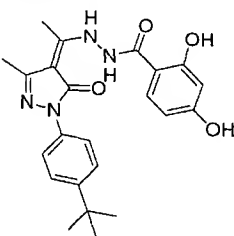
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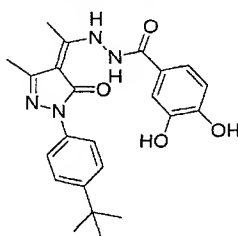
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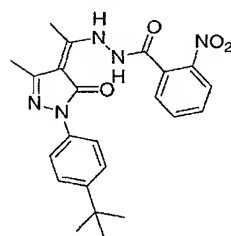
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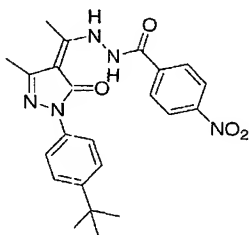
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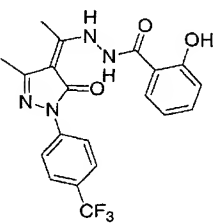
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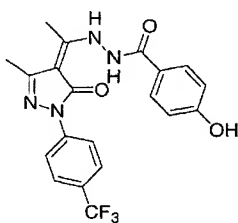
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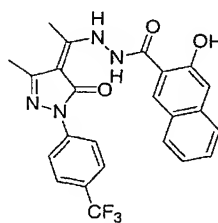
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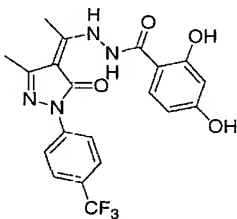
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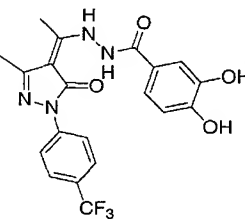
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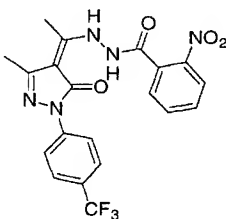
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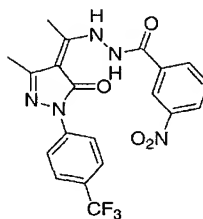
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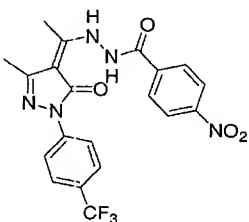
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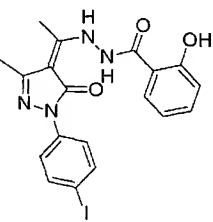
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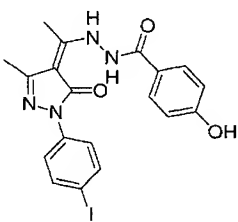
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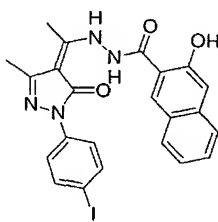
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SYNTHETIC EX. 39

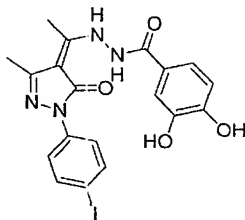


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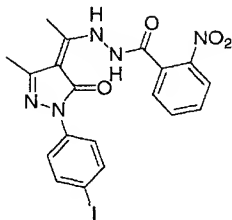


【Ka 20】

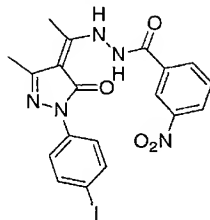
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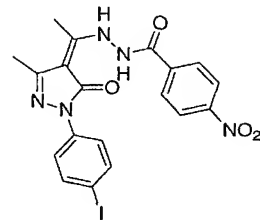
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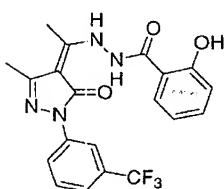
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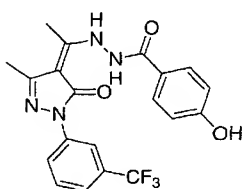
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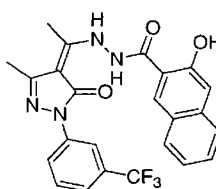
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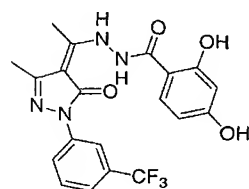
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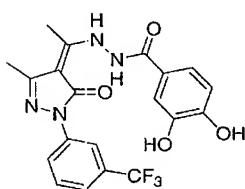
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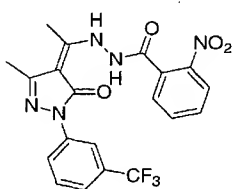
SYNTHETIC EX. 48



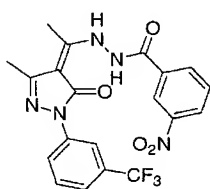
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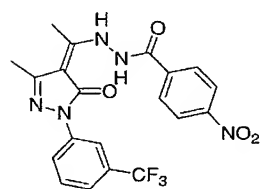
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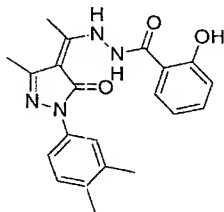
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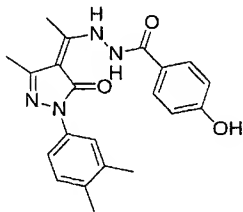
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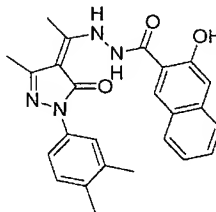
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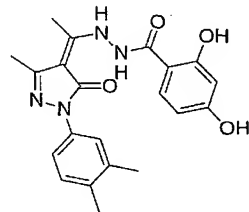
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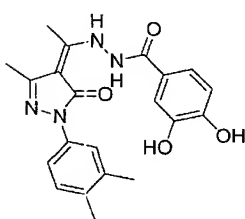
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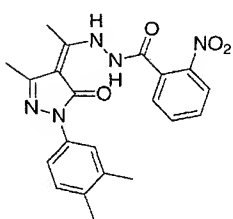
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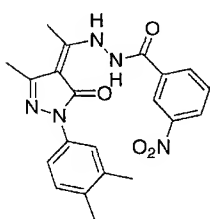
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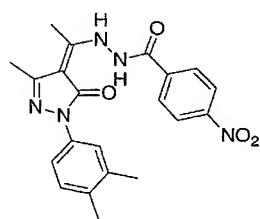
SYNTHETIC EX. 58



SYNTHETIC EX. 59

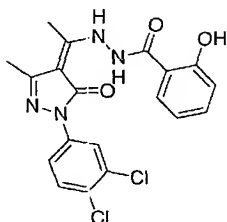


SYNTHETIC EX. 60

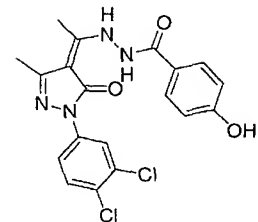


【Ka 21】

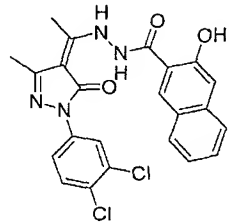
SYNTHETIC EX. 61



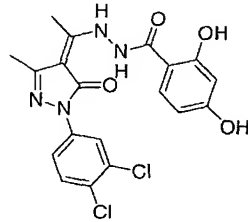
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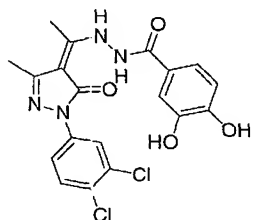
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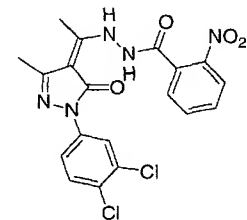
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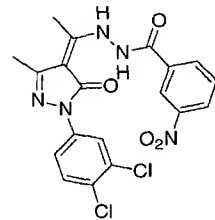
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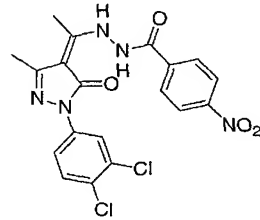
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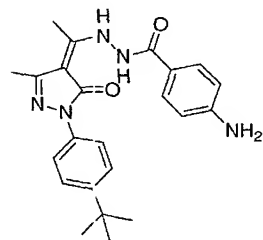
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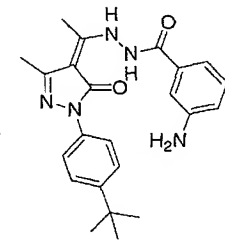
SYNTHETIC EX. 68



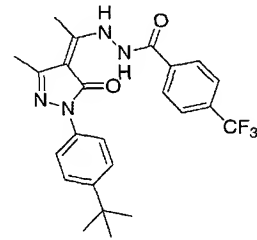
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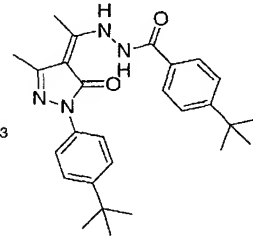
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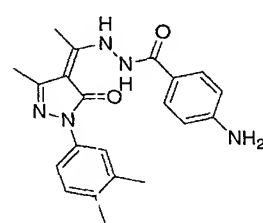
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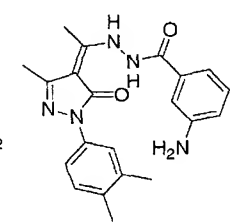
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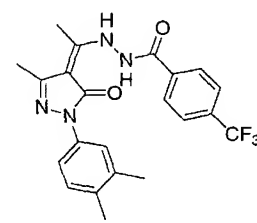
SYNTHETIC EX. 73



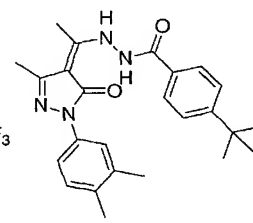
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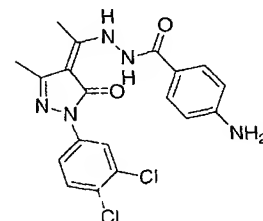
SYNTHETIC EX. 75



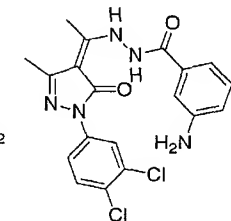
SYNTHETIC EX. 76



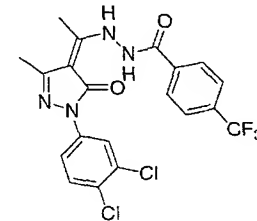
SYNTHETIC EX. 77



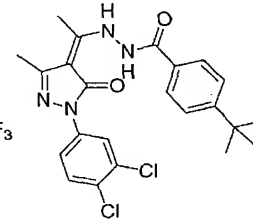
SYNTHETIC EX. 78



SYNTHETIC EX. 79

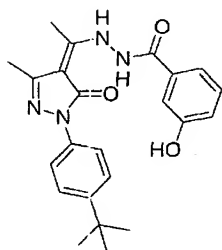


SYNTHETIC EX. 80

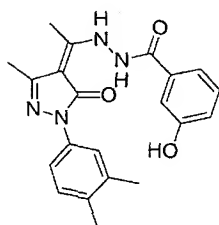


【Ka 22】

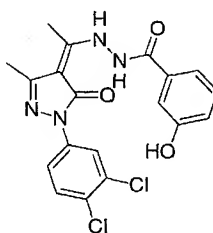
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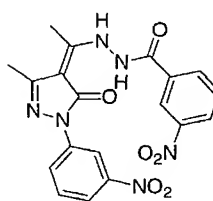
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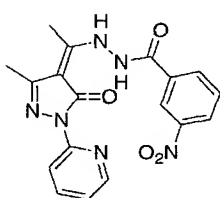
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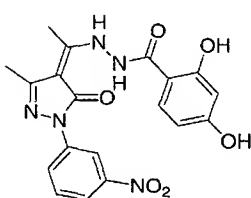
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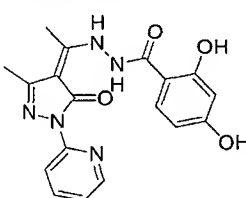
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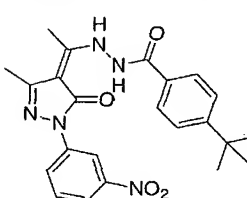
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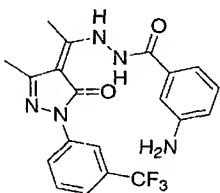
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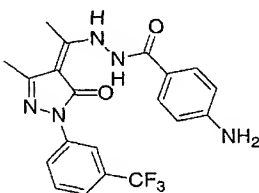
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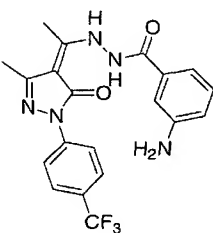
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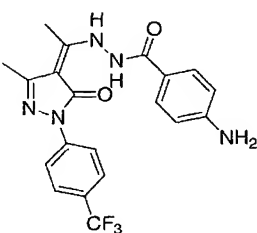
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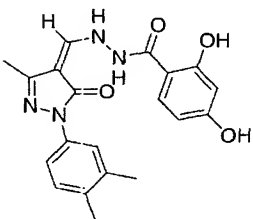
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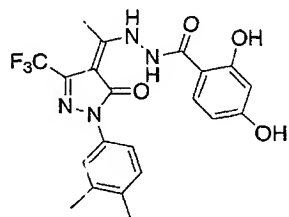


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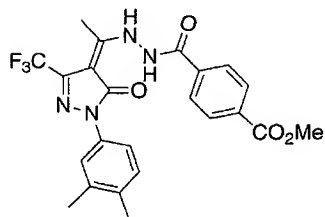


【Ka 23】

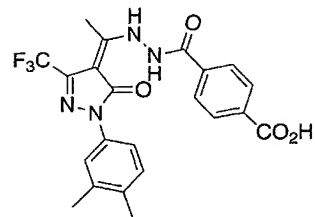
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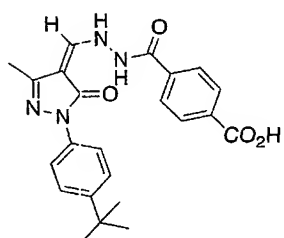
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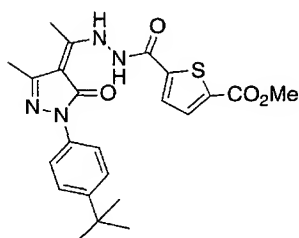
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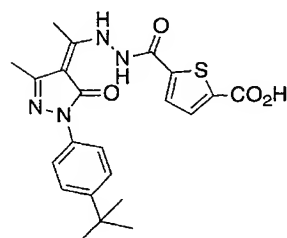
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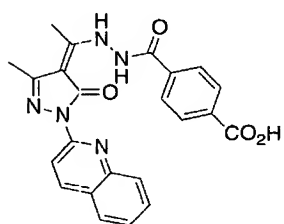
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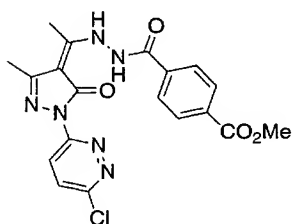
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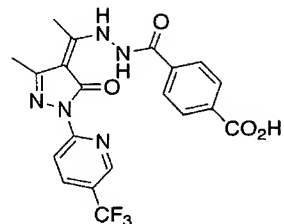
SYNTHETIC EX. 100



SYNTHETIC EX. 101



SYNTHETIC EX. 102



ASSAY EXAMPLE 1

Stimulation of Proliferation of a Thrombopoietin (TPO)-dependent Cell Line (1)

The reactivity of Synthetic Example 56, the compound
5 of the present invention, with thrombopoietin (TPO) receptor was assayed using a human leukemic cell line, UT7/EPO-*mpl*.

(1) Cells and cell culture

UT7/EPO-*mpl* is a stable transformed cell line
10 obtained by introducing into human leukemic cell line UT7/EPO a vector that induces expression of human thrombopoietin receptor (*c-mpl*) under control of a cytomegaloviral promoter by the method of Komatsu et al. (J. Biol. Chem., 272:7259-7263 (1997)). Proliferation of
15 this cell line is stimulated by thrombopoietin, while its mother cell line UT7/EPO exhibits no response to thrombopoietin. These two cell lines were subcultured in IMDM (GIBCO) containing 10% fetal bovine serum (TRACE SCIENTIFIC) using a CO2 incubator (5% CO2, 37°C).

20 (2) Cell proliferation assay by the MTT method

The subcultured cells described above were washed twice with PBS and suspended in IMDM containing 10% fetal bovine serum at a cell density of 6×10^4 cells/ml. The cell suspension was transferred to a 96-well tissue
25 culture plate (CORNING) in 100- μ l aliquots. Then Synthetic Example 56 dissolved in DMSO was diluted 100-fold with IMDM containing 10% fetal bovine serum and

added to the aforementioned cell suspension in 20- μ l aliquots. The suspension was incubated in a CO₂ incubator (5% CO₂, 37°C) for 4 days. Cell proliferation was assayed according to the method of Mosmann et al. (J. Immunological Methods, 65:55-63 (1983)). A 10- μ l aliquot of 5 mg/ml MTT reagent (SIGMA) was added to each well of the tissue culture plate and the plate was incubated at 37°C for 4 h. The formazan pigment generated was dissolved by adding 150 μ l per well of 0.1 M HCl/isopropanol solution, and the absorbance of the resulting pigment solution was measured at 550 nm with a 96-well microplate reader (BIO-RAD, M450). Figure 1 shows the results with UT7/EPO-mpl cells, while Figure 2 shows data obtained with UT7/EPO cells expressing no thrombopoietin receptor.

Figure 1 demonstrated that proliferation of UT7/EPO-mpl cells was stimulated by Synthetic Example 56 in a concentration-dependent manner, while no effect of this compound on proliferation was observed with UT7/EPO, the mother cell line, as shown in Figure 2.

ASSAY EXAMPLE 2

Activity of Signal Transduction Mediated by Thrombopoietin Receptor

The signal-transducing activity of Synthetic Example 56, the compound of the present invention, mediated by thrombopoietin receptor was assayed according to the method of Komatsu et al. (Blood, 87:4552-4560 (1996)).

Human leukemic cell line UT7/EPO-mpl was washed three times with PBS and suspended in IMDM (GIBCO) containing 10% fetal bovine serum (TRACE SCIENTIFIC) at a cell density of 9×10^5 cells/ml. The cell suspension was
5 incubated in a CO₂ incubator (5% CO₂, 37°C) for 18 h. To 2 ml of this cell suspension (7×10^6 cells/ml), either thrombopoietin (final concentration, 30 ng/ml) or a DMSO solution of Synthetic Example 56 (final concentration, 1 µg/ml) was added. After incubating the mixture at 37°C
10 for 1-15 min, the cells were lysed in 1.4 ml of TNE buffer [20 mM Tris-HCl buffer (pH 7.4) containing 150 mM NaCl, 1 mM EDTA, 1% Triton X-100, 1 mM PMSF, 1 mM Na₃VO₄, and 1/400-diluted Protease inhibitor cocktail (SIGMA)]. The cell lysate was centrifuged to collect the
15 supernatant for immunoprecipitation with antibodies against proteins involved in signal transduction [anti-STAT3 (SANTACRUZ BIOTECHNOLOGY) and anti-STAT5A (UPSTATE BIOTECHNOLOGY)] and protein G Sepharose (PHARMACIA). The immunoprecipitated protein fraction was collected and
20 denatured in a sample buffer for separation by SDS-polyacrylamide gel electrophoresis (7.5%). The separated proteins were transferred onto PVDF membrane (ATTO, 0.2 µm) at 100 V for 1 h for detection of tyrosine phosphorylation using an alkaline phosphatase-labelled
25 antibody against phosphorylated tyrosine (RC20, TRANSDUCTION LABORATORIES). The antigen-antibody complex formed on the PVDF membrane was visualized with 150 µg/ml

NBT (BIO-RAD) and 300 µg/ml BCIP (BIO-RAD). The results are summarized in Table 7.

Table 7

	DMSO	SYNTHETIC EXAMPLE No. 56	Thrombopoietin
STAT 3	-	+	+
STAT 5A	-	+	+

5 ASSAY EXAMPLE 3

The following Synthetic Examples were tested according to the method of Assay Example 1 to determine the maximal growth rate (Efficacy), expressed by taking the value with human leukemic cell line UT7/EPO-mpl
10 observed in the presence of 10 ng TPO as 100% standard, and the concentration of each compound that yields a growth rate corresponding to 50% of the maximum cell growth observed with the same compound (EC₅₀). The results are summarized in Table 8. (Here, "-" indicates
15 that EC₅₀ was not determined because the value of Efficacy was below the detection limit.)

Table 8

Synthetic Example No.	Efficacy (%)	EC ₅₀ (ng/ml)
1	74	7.4
2	89	6.3
3	82	15
4	53	15
5	86	3.4
6	64	7.4
7	99	2.2
8	52	31
9	90	5.1
10	78	20
11	83	2.0
12	100	76
13	99	280
14	91	72
15	109	23
16	58	61
17	73	79
18	94	55
19	100	14
20	91	38
21	39	290
22	50	190
23	129	28
24	89	7.2
25	54	200
26	78	2.9
27	75	5.6
28	99	37
29	67	230
30	106	19
31	63	5.2
32	90	37
33	96	1.1
34	99	5.2
35	99	34
36	97	59
37	63	140
38	93	36
39	97	28
40	37	250
41	115	32
42	71	250
43	87	83
44	26	250

45	74	30
46	82	15
47	48	190
48	62	8.0
49	62	9.1
50	89	37
51	73	33
52	22	120
53	120	12
54	61	7.5
55	53	220
56	96	1.1
57	97	5.9
58	110	32
59	82	24
60	62	100
61	91	29
62	57	6.4
63	21	190
64	74	7.7
65	70	8.9
66	133	33
67	80	33
68	26	210
69	89	5.7
70	87	23
71	89	69
72	88	75
73	84	10
74	77	25
75	89	63
76	79	46
77	78	5.1
78	69	15
79	81	160
80	71	640
81	84	7.2
82	84	26
83	78	6.1
84	109	130
86	105	21
87	71	600
88	70	130
89	68	39
90	76	21
91	81	24
92	82	5.5
93	84	4.3

Reference		
Synthetic Example	7	-
1		
Reference		
Synthetic Example	12	-
2		
Reference		
Synthetic Example	7	-
3		
Reference		
Synthetic Example	67	1400
4		

ASSAY EXAMPLE 4

Synthetic Example 56, the compound of the present invention, and four compounds (Reference Synthetic
5 Examples 1 to 4) described in a publication of international patent application, Publication No. WO01/34585, applied by SmithKlineBeecham Corp were tested according to the method of Assay Example 1. Figure 3 shows the results.

10 ASSAY EXAMPLE 5

Activity of Stimulating Proliferation of a Thrombopoietin (TPO)-dependent Cell Line (2)

Human leukemic cell line UT7/EPO-mpl was washed twice with PBS and suspended in IMDM containing 10% fetal
15 bovine serum at a cell density of 6×10^4 cells/ml. The cell suspension was transferred to a 96-well tissue culture plate (CORNING) in 100- μ l aliquots. Then the following Synthetic Examples, each dissolved in DMSO, were diluted 83-fold with IMDM containing 10% fetal
20 bovine serum and added to the aforementioned cell

suspension in 20- μ l aliquots. The suspension was incubated in a CO₂ incubator (5% CO₂, 37°C) for 4 days. Cell proliferation was assayed using WST-8 reagent (Kishida Chemical, Co. Ltd.) according to instructions by the manufacturer. A 10- μ l aliquot of 5 mM WST-8 reagent solution was added to each well of the tissue culture plate, and the plate was incubated at 37°C for 4 h. The formazan pigment generated was detected by measuring the absorbance at 450 nm with a 96-well microplate reader (Nihon Molecular Devices, Spectramax 190). The concentration of each compound that yields a growth rate corresponding to 50% of the growth of human leukemic cell line UT7/EPO-mpl observed in the presence of 10 ng/ml TPO (EC₅₀T) and the maximal growth rate achieved by the same compound (Efficacy), expressed by taking the value with human leukemic cell line UT7/EPO-mpl in the presence of 10 ng/ml TPO as 100% standard, are summarized in Table 9.

Table 9

Synthetic Example No.	Efficacy (%)	EC ₅₀ T (ng/ml)
94	95	3.3
95	71	52
96	93	3.3
97	94	25
98	96	31
99	110	3.9
100	107	59
101	100	18
102	97	69

FORMULATION EXAMPLE 1

A granule preparation containing the following

ingredients is prepared.

Ingredients

Compound represented by the formula (1)	10 mg
Lactose	700 mg
Corn Starch	274 mg
HPC-L	16 mg
	1000 mg

A compound represented by the formula (1) and lactose are sifted through a 60-mesh sieve. Corn starch is
5 sifted through a 120-mesh sieve. They are mixed in a V-type blender. The powder mixture is kneaded with a low-viscosity hydroxypropylcellulose (HPC-L) aqueous solution, granulated (extrusion granulation, die size 0.5-1 mm) and dried. The resulting dry granules are sifted through a
10 shaking sieve (12/60 mesh) to obtain a granule preparation.

FORMULATION EXAMPLE 2

A powder preparation for capsulation containing the following ingredients is prepared.

15 Ingredients

Compound represented by the formula (1)	10 mg
Lactose	79 mg
Corn Starch	10 mg
Magnesium Stearate	1 mg
	100 mg

A compound represented by the formula (1) and lactose are sifted through a 60-mesh sieve. Corn starch is sifted through a 120-mesh sieve. They are mixed with magnesium stearate in a V-type blender. The 10% powder
20 is put in hard capsules No. 5, 100 mg each.

FORMULATION EXAMPLE 3

A granule preparation for capsulation containing the following ingredients is prepared.

Ingredients

Compound represented by the formula (1)	15 mg
Lactose	90 mg
Corn Starch	42 mg
HPC-L	3 mg
	<hr/> 150 mg

- 5 A compound represented by the formula (1) and lactose are sifted through a 60-mesh sieve. Corn starch is sifted through a 120-mesh sieve. They are mixed in a V-type blender. The powder mixture is kneaded with a low-viscosity hydroxypropylcellulose (HPC-L) aqueous solution,
- 10 granulated and dried. The resulting dry granules are sifted through a shaking sieve (12/60 mesh). The granules are put in hard capsules No. 4, 150 mg each.

FORMULATION EXAMPLE 4

- 15 A tablet preparation containing the following ingredients is prepared.

Ingredients

Compound represented by the formula (1)	10 mg
Lactose	90 mg
Microcrystalline cellulose	30 mg
Magnesium Stearate	5 mg
CMC-Na	15 mg
	<hr/> 150 mg

- 20 A compound represented by the formula (1), lactose, microcrystalline cellulose and CMC-Na (carboxymethylcellulose sodium salt) are sifted through a 60-mesh sieve and mixed. The powder mixture is mixed with magnesium stearate to give a bulk powder mixture.

The powder mixture is compressed directly into 150 mg tablets.

FORMULATION EXAMPLE 5

An intravenous preparation is prepared as follows.

Compound represented by the formula (1)	100 mg
Saturated Fatty Acid Glyceride	1000 ml

- 5 Solutions having the above-mentioned composition are usually administered to a patient intravenously at a rate of 1 ml per 1 minute.

【Industrial applicability】

- 10 The compounds of the present invention which have affinity for thrombopoietin receptor and act as thrombopoietin receptor agonists are useful as preventive, therapeutic and improving agents for diseases against which activation of the thrombopoietin receptor is effective, especially as drugs for hematological
- 15 disorders accompanied by abnormal platelet count and as drugs for diseases treated or prevented by stimulating differentiation and proliferation of vascular endothelial cells and endothelial progenitor cells, and are useful as medicines.

- 20 【Brief description of drawings】

【Fig. 1】

The proliferation of UT7/EPO-mpl cells when stimulated by a compound of the present invention (Synthetic Example 56) assayed by the MTT method.

- 25 【Fig. 2】

The proliferation of UT7/EPO cells when stimulated

by a compound of the present invention (Synthetic Example 56) assayed by the MTT method.

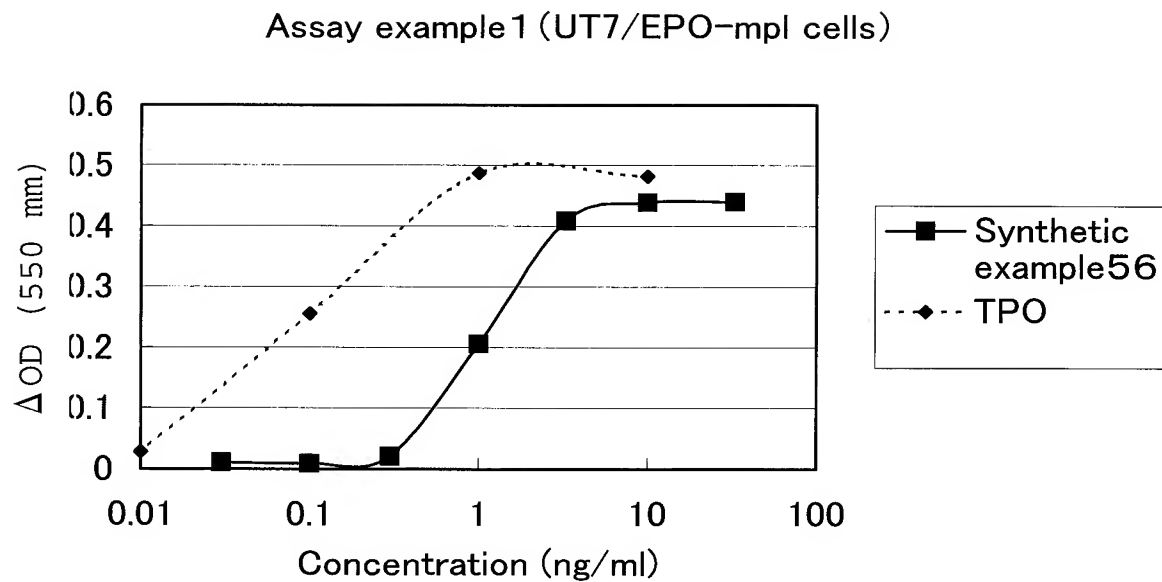
【Fig. 3】

The proliferation of UT7/EPO-mpl cells when
5 stimulated by a compound of the present invention
(Synthetic Example 56) or the compounds described in a
publication of international patent application
(Reference Synthetic Examples 1 to 4) assayed by the MTT
method is shown.

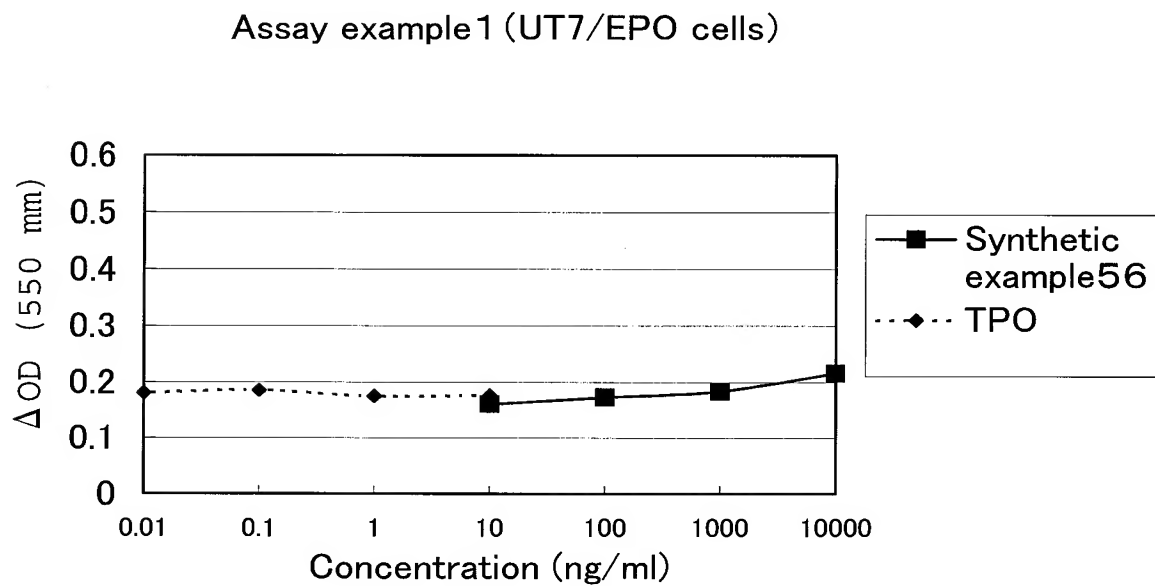
【Type of Document】

DRAWING

【Fig. 1】

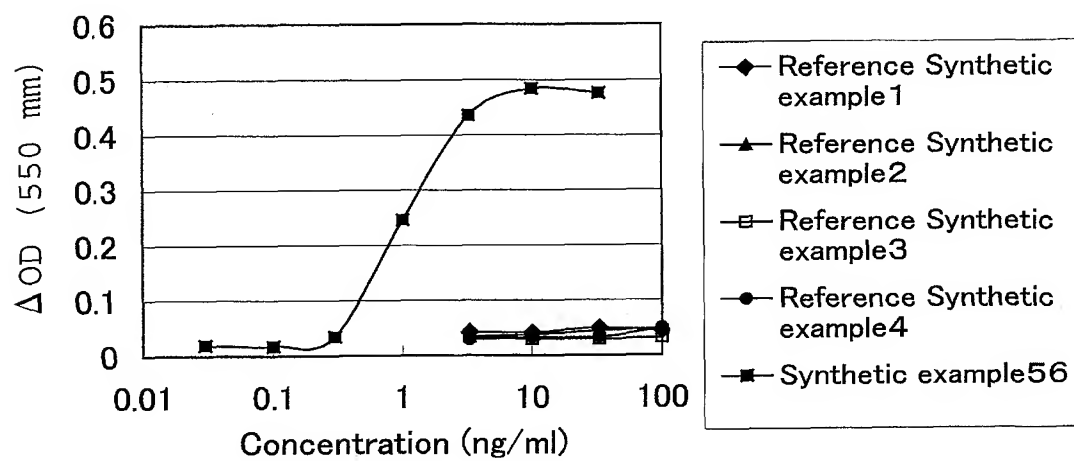


【Fig. 2】



【Fig. 3】

Assay example4 (UT7/EPO-mpl cells)



【Type of Document】

ABSTRACT

【Summary】

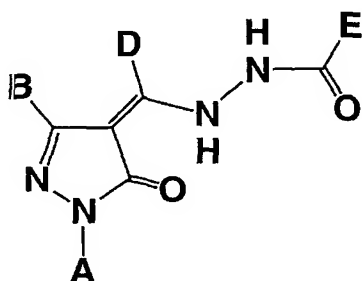
【Object】

To provide a preventive, therapeutic or improving
5 agent for diseases against which activation of the
thrombopoietin receptor is effective.

【Means for solving problem】

A preventive, therapeutic or improving agent for
diseases against which activation of the thrombopoietin
10 receptor is effective or a platelet increasing agent,
which contains a thrombopoietin receptor activator
represented by the formula (1):

【Ka 1】



15 [wherein A is a C₂₋₁₄ aryl group, B is hydrogen, a C₁₋₆
alkyl group, a C₁₋₃ alkyl group substituted with one or
more fluorine atoms or a C₂₋₁₄ aryl group, D is hydrogen,
a C₁₋₆ alkyl group, a C₁₋₃ alkyl group substituted with one
or more fluorine atoms or a C₂₋₁₄ aryl group, and E is a
20 C₂₋₁₄ aryl group], a tautomer, prodrug or pharmaceutically
acceptable salt of the activator or a solvate thereof, as
an active ingredient.

【Selected Figure】

No Selected Figure